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U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

**TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371**

ATTORNEY'S DOCKET NUMBER

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U.S. APPLICATION NO.
(If known, see 37CFR1.5)**10/030481**

INTERNATIONAL APPLICATION NO.

PCT/EP00/06788

INTERNATIONAL FILING DATE

July 17, 2000

PRIORITY DATE CLAIMED

July 23, 1999

TITLE OF INVENTION

COMPOUND

APPLICANT(S) FOR DO/EO/US

Guy MARCK and Andreas SCHUSTER

Applicant(s) herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.
4. ☒ The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c)(2)).
 - a. ☒ is attached hereto (required only if not communicated by the International Bureau.
 - b. ☒ has been communicated by the International Bureau.
 - c. ☐ is not required, as the application was filed with the United States Receiving Office (RO/US).
6. ☐ An English language translation of the International Application as filed (35 U.S.C. 371 (c)(2)).
 - a. ☐ is attached hereto.
 - b. ☐ has been previously submitted under 35 U.S.C. 154 (d)(4).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3)).
 - a. ☐ are attached hereto (required only if not communicated by the International Bureau).
 - b. ☐ have been communicated by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371 (c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
10. ☐ An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).

Items 11 to 20 below concern document(s) or information included:

11. ☒ Information Disclosure Statement under 37 CFR 1.97 and 1.98
12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A **FIRST** preliminary amendment, with Appendix and Abstract of the Disclosure.
14. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
15. ☐ A Substitute specification.
16. ☐ A change of power of attorney and/or address letter.
17. ☐ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821-1.825.
18. ☐ A second copy of the published international application under 35 U.S.C. 154 (d)(4).
19. ☐ A second copy of the English language translation of the international application 35 U.S.C. 154 (d)(4).
20. ☒ Other items or information:
 - a. ☒ Copy of cover page of International Publication No. WO01/07495 A1.
 - b. ☐ Copy of Notification of Missing Requirements.
 - c. ☐

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:)
)
Guy Marck et al.) Group Art Unit: Unassigned
)
Application No.: Unassigned) Examiner: Unassigned
)
Int'l. Filing Date: July 17, 2000)

For: COMPOUND, being a National Stage filing of PCT International Application No. PCT/EP00/06788, filed on July 17, 2000

Assistant Commissioner for Patents
Washington, DC 20231

Sir:

Preliminary Amendment

Before examining this application, please amend the application as follows:

In the specification:

Page 1, after the title, insert as a first paragraph the following:

-- This application is a national stage filing under 35 U.S.C. § 371 of international application no. PCT/EP00/06788, filed on July 17, 2000, which published in the English language. This application also claims the benefit of priority under 35 U.S.C. § 119(a) to European patent application no. 99305857.7, filed on July 23, 1999. --

In the claims:

Please cancel claims 8-9 without prejudice or disclaimer

Please also amend claims 3-7 and 10-13 as follows:

3. (Amended) A compound according to Claim 1, in which the repeating unit of formula (I) accounts for at least 50% of the monomer building blocks in the compound.

4. (Amended) A compound according to claim 1, in which the group M is selected from acrylate; methacrylate; 2-chloroacrylate; 2-phenylacrylate; acrylamide, methacrylamide, 2-chloroacrylamide and 2-phenylacrylamide, the nitrogen atom of which is optionally substituted by a lower alkyl group; vinyl ether; vinyl ester; a styrene derivative; siloxane; imide; amic acid; an amic acid ester; amidimide; a maleic acid derivative and a fumaric acid derivative.

5. (Amended) A method of manufacturing a compound as claimed in claim 1, comprising the polymerization of one or more pre-finished monomer units of formula (I).

6. (Amended) A method of manufacturing a compound as claimed in claim 1, which comprises reacting a photoactive derivative with a functional polymer analogue of a polymer according to Claim 1.

7. (Amended) A polymer layer, comprising a compound of formula (I) in cross-linked form.

10. (Amended) An optical or an electro-optical device, comprising a compound according to claim 1.

11. (Amended) An optical or an electro-optical device, comprising a layer according to Claim 7.

12. (Amended) A compound as claimed in claim 1, which is Poly-[1-[11-[5-[4-[(E)-2-methoxy-carbonylvinyl]benzoyloxy]-2-[6-[2-methoxy-(E)-4-(methoxycarbonylvinyl)-phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene].

13. (Amended) A compound as claimed in claim 1, which is Poly-[1-[11-(*E,E*)-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxy]undecyloxy]-1-methylethylene].

In the abstract:


Please add the abstract attached to the end of this Preliminary Amendment.

Remarks

Claims 1-7 and 10-13 are pending in this application. All claims have been amended for only formal and stylistic reasons, and none of the amendments change the scope of the claims. If there is any fee due in connection with the filing of this Preliminary Amendment, please charge the fee to our Deposit Account No. 06-0916.

Respectfully submitted,

FINNEGAN, HENDERSON, FARABOW,
GARRETT & DUNNER, L.L.P.

By: 
Steven J. Scott
Reg. No. 43,911

Date: January 7, 2002

Appendix Detailing Amendments to Claims

3. (Amended) A compound according to Claim 1 [or Claim 2], in which the repeating unit of formula (I) accounts for [comprises] at least 50% of the monomer building blocks in [comprising] the compound [of formula (I)].

4. (Amended) A compound according to claim 1 [any one of claims 1 to 3], in which the group M is selected from acrylate; methacrylate; 2-chloroacrylate; 2-phenylacrylate; acrylamide, methacrylamide, 2-chloroacrylamide and 2-phenylacrylamide, the nitrogen atom of which is optionally substituted by a lower alkyl group; vinyl ether; vinyl ester; a styrene derivative; siloxane; imide; amic acid; an amic acid ester [esters]; amidimide; a maleic acid derivative [derivatives] and a fumaric acid derivative [derivatives].

5. (Amended) A method of manufacturing a compound as claimed in claim 1 [of formula (I)], comprising the polymerization of one or more pre-finished monomer units of formula (I).

6. (Amended) A method of manufacturing a compound as claimed in claim 1, which comprises [of formula (I) comprising] reacting a photoactive derivative with a functional polymer analogue of a polymer according to Claim 1.

7. (Amended) A polymer layer₁ comprising a compound of formula (I) in cross-linked form.

10. (Amended) An optical or an electro-optical device₁ comprising a compound according to claim 1 [any one of claims 1 to 4].

11. (Amended) An optical or an electro-optical device₁ comprising a layer according to Claim 7 [or Claim 8].

12. (Amended) A compound as claimed in claim 1 [of formula (I)], which is Poly-[1-[11-[5-[4-[(*E*)-2-methoxy-carbonylvinyl]benzoyloxy]-2-[6-[2-methoxy-(*E*)-4-(methoxycarbonylvinyl)-phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene].

13. (Amended) A compound as claimed in claim 1 [of formula (I)], which is Poly-[1-[11-(*E,E*)-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene].

10/030481
531 Rec'd PCT/EP 09 JAN 2002Compound

The present invention relates to new photoactive polymers, their use as liquid crystal (LC) orientation layers and their use in the construction of unstructured and structured optical and electro-optical elements and multi-layer systems.

5 The successful functioning of a Liquid Crystal Device relies upon the ability of the LC molecules within that device to assume and maintain an alignment imposed upon them. Alignment of the LC molecules is achieved by use of an orientation layer which defines a direction of orientation for the LC molecules of the device with the result that the longitudinal axes of the molecules become aligned with the direction of
10 orientation defined by the orientation layer. In addition to this directional alignment, the orientation layer is also able to impart to the LC molecules an angle of tilt so that the molecules align themselves at an angle to the surface of the orientation layer rather than lying parallel thereto.

15 Tilt angles of between 1° and 15° are usual for Nematic LCDs. Tilt angles of about 7° are required for supertwisted nematic (STN) LCDs in order to avoid the formation of so-called fingerprint textures. Vertically aligned nematic (VAN) LCD's for instance require pretilt angles of between 85° and 90° .

20 Methods of preparing structured and unstructured orientation layers are well known to a skilled person. In particular it is known that by using linearly polarised light it is possible to prepare orientation layers in which both the direction of orientation and the tilt angle of the orientation layer are determined by the direction and angle of incidence of the plane polarised light used to irradiate said layer.

25 Structured orientation layers are of great interest in many areas of display technology and integrated optics. These layers are characterised by regions (pixels) which alternate in respect of the direction of orientation and angle of tilt of their component molecules. These orientation layers can be used to improve the viewing angle dependency of TN, STN and VAN LCDs, for example:

A possible method of producing high-resolution structured orientation patterns in liquid crystalline layers is described in *Jpn. J. Appl. Phys.*, Vol. 31 (1992), 2155. In

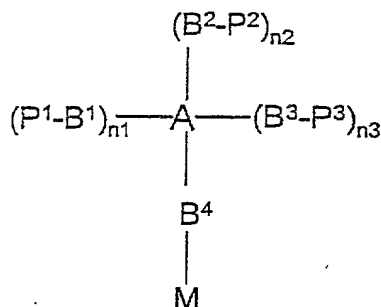
that process the dimerisation of polymer-bonded photoreactive cinnamic acid groups induced by irradiation with linearly polarised light is employed for the structured orientation of liquid crystals. Those photo-oriented polymer networks can be used wherever structured or unstructured liquid crystal orientation layers are required. In addition to their use in LCDs, these orientation layers can also be used, for example, in the production of so-called hybrid layers, as illustrated in European Patent Applications EP-A-0 611 981, EP-A-0 689 084 and EP-A-0 689 065. It is possible, using these hybrid layers of photostructured orientation polymers and crosslinkable low molecular weight liquid crystals, to prepare optical elements such as, non-absorptive colour filters, linear and circular polarisers, optical delay layers and so on.

The ability of the resulting orientation layers to perform their function thus depends, in part, upon the number of molecules in the layer that have been dimerised as a result of irradiation with linearly polarised light. The extent to which the molecules are dimerised relies, in part, on the irradiation time, the irradiation energy and the structure of the molecules being irradiated.

EP-A-0 611 786, EP-A-0 763 552, EP-A-0 860 455, WO 96/10049 and WO 99/15576 describe polymers that are suitable in principle for the production of such anisotropically crosslinked, photostructured orientation layers for liquid crystals.

However, a problem with many polymers currently used in the preparation of photo-orientated orientation layers is that relatively long irradiation times are required to effect efficient dimerisation of the component molecules. There is, therefore, a need for photo crosslinkable polymers that can be readily cross-linked over a relatively short irradiation time. The present invention addresses that need.

A first aspect of the present invention provides a polymeric compound comprising a repeating unit of formula (I)



I

in which:

5 A represents a nitrogen atom, a carbon atom, a group $-CR^1-$ or an aromatic or alicyclic group, which is optionally substituted by a group selected from fluorine, chlorine, cyano and a C_{1-18} cyclic, straight-chain or branched alkyl group, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkyl

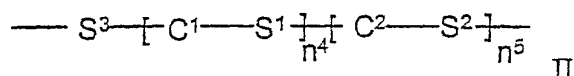
10 $-CH_2-$ groups are optionally replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-Si(CH_3)_2-O-Si(CH_3)_2-$, $-NR^1-$, $-NR^1-CO-$, $-CO-NR^1-$, $-NR^1-CO-O-$, $-O-CO-NR^1-$, $-NR^1-CO-NR^1-$, $-CH=CH-$, $-C\equiv C-$ and $-O-CO-O-$, wherein R^1 represents a hydrogen atom or lower alkyl,

15 M represents a repeating monomer unit;

n^1 to n^3 each independently represent 0 or an integer having a value of from 1 to 3, with the proviso that $1 < n^1 + n^2 + n^3 < 4$;

P^1, P^2, P^3 each independently represents a photoactive group; and

B^1 to B^4 each independently represent a residue of general formula II



in which

S¹ to S³ each independently represent a single bond or a spacer group selected from a C₁₋₂₄ straight-chain or branched alkylene group, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkylene -CH₂- groups are optionally replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -Si(CH₃)₂-O-Si(CH₃)₂-, -NR¹-, -NR¹-CO-, -CO-NR¹-, -NR¹-CO-O-, -O-CO-NR¹-, -NR¹-CO-NR¹-, -CH=CH-, -C≡C- and -O-CO-O- wherein R¹ is as defined above,

C¹ and C² each independently represents an aromatic or an alicyclic group, which is optionally substituted by a group selected from fluorine, chlorine, cyano or a C₁₋₁₈ cyclic, straight-chain or branched alkyl group, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkyl -CH₂- groups are optionally replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -Si(CH₃)₂-O-Si(CH₃)₂-, -NR¹-, -NR¹-CO-, -CO-NR¹-, -NR¹-CO-O-, -O-CO-NR¹-, -NR¹-CO-NR¹-, -CH=CH-, -C≡C- and -O-CO-O- wherein R¹ represents a hydrogen atom or lower alkyl, and

n⁴ and n⁵ are each independently 0 or 1.

The polymeric compounds of the present invention can be readily aligned upon exposure to linearly polarised light. In addition, by using the compounds of the invention, it is possible to reduce the irradiation time required to form cross-linked polymer films.

By the term "aromatic" it should be understood to include optionally substituted carbocyclic and heterocyclic groups.

By the term "cyclic, straight-chain or branched alkyl group, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent $-\text{CH}_2-$ groups are optionally replaced by a group selected from $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-$, $-\text{CH}=\text{CH}-$ and $-\text{C}\equiv\text{C}-$," it should be understood to include groups selected from the group comprising methyl, ethyl, propyl, isopropyl, butyl, isobutyl, *sec*-butyl, *tert*-butyl, pentyl, isopentyl, cyclopentyl, hexyl, cyclohexyl, heptyl, octyl, nonyl, decyl, undecyl, dodecyl, 3-methylpentyl, allyl, but-3-en-1-yl, pent-4-en-1-yl, hex-5-en-1-yl, propynyl, butynyl, pentynyl, methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, *sec*-butoxy, *tert*-butoxy, pentyloxy, isopentyloxy, cyclopentyloxy, hexyloxy, cyclohexyloxy, heptyloxy, octyloxy, nonyloxy, decyloxy, undecyloxy, dodecyloxy, 3-methylpentyloxy, allyloxy, but-3-enyloxy, pent-4-enyloxy, cyclohexylmethoxy, cyclopentylmethoxy, methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, butoxycarbonyl, isobutoxycarbonyl, *sec*-butoxycarbonyl, *tert*-butoxycarbonyl, pentyloxycarbonyl, isopentyloxycarbonyl, cyclopentyloxycarbonyl, hexyloxycarbonyl, cyclohexyloxycarbonyl, octyloxycarbonyl, nonyloxycarbonyl, decyloxycarbonyl, undecyloxycarbonyl, dodecyloxycarbonyl, 3-methylpentyloxycarbonyl, allyloxycarbonyl, but-3-enyloxycarbonyl, pent-4-enyloxycarbonyl, cyclohexylmethoxycarbonyl, cyclopentylmethoxycarbonyl, acetoxycarbonyl, ethylcarbonyloxy, propylcarbonyloxy, isopropylcarbonyloxy, butylcarbonyloxy, isobutylcarbonyloxy, *sec*-butylcarbonyloxy, *tert*-butylcarbonyloxy, pentylcarbonyloxy, isopentylcarbonyloxy, cyclopentylcarbonyloxy, hexylcarbonyloxy, cyclohexylcarbonyloxy, octylcarbonyloxy, nonylcarbonyloxy, decylcarbonyloxy, undecylcarbonyloxy, dodecylcarbonyloxy, 3-methylpentylcarbonyloxy, but-3-enyloxy, pent-4-enyloxy, acetyl, ethylcarbonyl, propylcarbonyl, isopropylcarbonyl, butylcarbonyl, isobutylcarbonyl, *sec*-butylcarbonyl, pentylcarbonyl, isopentylcarbonyl, cyclohexylcarbonyl, octylcarbonyl, nonylcarbonyl, decylcarbonyl, undecylcarbonyl, dodecylcarbonyl, methoxyacetoxycarbonyl,

1-methoxy-2-propoxy, 3-methoxy-1-propoxy, 2-methoxyethoxy, 2-isopropoxyethoxy, 1-ethoxy-3-pentyloxy, 3-butyloxy, 4-pentyloxy, 5-chloropentynyl, 4-pentyne-carbonyloxy, 6-propyloxyhexyl, 6-propyloxyhexyloxy, 2-fluoroethyl, trifluoromethyl, 2,2,2-trifluoroethyl, 1*H*,1*H*-pentadecafluorooctyl, 1*H*,1*H*,7*H*-dodeca-
 5 fluoroheptyl, 2-(perfluorooctyl)ethyl, 2-(perfluorobutyl)ethyl, 2-(perfluorohexyl)ethyl, 2-(perfluorodecyl)ethyl, perfluoropropyl, perfluorobutyl, perfluoroheptyl, perfluorooctyl, perfluorononyl. 1-fluoropropoxy, 1-fluoropentyloxy, 2-fluoropropoxy, 2,2-difluoropropoxy, 3-fluoropropoxy, 3,3-difluoropropoxy, 3,3,3-trifluoropropoxy, trifluoromethoxy and the like.

10 By the term "lower alkyl" it should be understood to include straight chain and branched hydrocarbon radicals having from 1 to 6 carbon atoms, preferably from 1 to 3 carbon atoms. Methyl, ethyl, propyl and isopropyl groups are especially preferred.

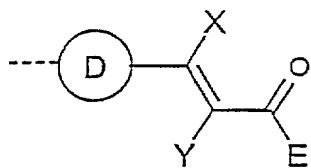
By the term "alicyclic" it should be understood to include non-aromatic carbocyclic or heterocyclic ring systems with 3 to 20 carbon atoms.

15 The group A is preferably an optionally substituted aromatic group. It is also preferred that when the group A is optionally substituted by an alkyl group, one or more of the alkyl -CH₂- groups are optionally replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO- and -CH=CH-.

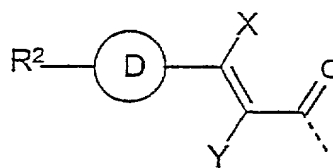
20 It is especially preferred that A is selected from the group selected from 1,2,5-benzenetriyl, 1,3,5-benzenetriyl and 1,3,4,5-benzenetetrayl, which are optionally substituted by one or more fluorine atoms. The group B⁴ preferably occupies position 1 of the especially preferred group A.

25 It is preferred that when $n^1 + n^2 = 2$ and $n^3 = 0$, A represents -CR¹- or an optionally substituted aromatic or alicyclic group. Alternatively, when $n^1 = 3$ and $n^2 + n^3 = 0$, A represents an optionally substituted aromatic or alicyclic group only.

The groups P¹, P² and P³ can be photoisomerised or photodimerised on exposure to UV or laser light. The groups P¹ to P³ preferably undergo photocyclisation reactions. The groups P¹ to P³ are preferably represented by the general formulae IIIa and IIIb:



IIIa



IIIb

5

wherein the broken line indicates the point of linkage to S^3 and wherein:

D represents pyrimidine-2,5-diyl, pyridine-2,5-diyl, 2,5-thiophenylene, 2,5-furanylene, 1,4- or 2,6-naphthylene; a phenylene group, which is optionally substituted by a group selected from fluorine, chlorine, cyano; or a C_{1-18} cyclic, straight-chain or branched alkyl residue, which is optionally substituted by a single cyano group or by one or more halogen groups and in which one or more non-adjacent alkyl $-CH_2-$ groups are optionally replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-Si(CH_3)_2-O-Si(CH_3)_2-$, $-NR^1-$, $-NR^1-CO-$, $-CO-NR^1-$, $-NR^1-CO-O-$, $-O-CO-NR^1-$, $-NR^1-CO-NR^1-$, $-CH=CH-$, $-C\equiv C-$ and $-O-CO-O-$, wherein R^1 is as defined above;

E represents $-OR^3$, $-NR^4R^5$ or an oxygen atom, which defines together with the ring D a coumarin unit, wherein R^3 , R^4 and R^5 are selected from hydrogen and a C_{1-18} cyclic, straight-chain or branched alkyl residue, which is optionally substituted by one or more halogen atoms and in which one or more non-adjacent alkyl $-CH_2-$ groups are optionally replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$ and $-CH=CH-$, or R^4 and R^5 together form a C_{5-8} alicyclic ring;

X, Y each independently represent hydrogen, fluorine, chlorine, cyano or a C₁₋₁₂ alkyl group, which is optionally substituted by fluorine and in which one or more non-adjacent alkyl -CH₂- groups are optionally replaced by a group selected from -O-, -CO-O-, -O-CO- and -CH=CH-;

5 R² represents hydrogen or a C₁₋₁₈ straight-chain or branched alkyl residue, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkyl -CH₂- groups are independently optionally replaced by a group selected from -O-, -CO-,
 10 -CO-O-, -O-CO-, -Si(CH₃)₂-O-Si(CH₃)₂-, -NR¹-, -NR¹-CO-,
 -CO-NR¹-, -NR¹-CO-O-, -O-CO-NR¹-, -NR¹-CO-NR¹-,
 -CH=CH-, -C≡C- and -O-CO-O-, wherein R¹ is as defined above.

It is preferred that the groups X and Y represent hydrogen.

It is also preferred that the group D is selected from pyrimidine-2,5-diyl, pyridine-2,5-diyl, 2,5-thiophenylene, 2,5-furanylene, 1,4- or 2,6-naphthylene and a
 15 phenylene group, which is optionally substituted by a C₁₋₁₂ cyclic, straight-chain or branched alkyl residue, which alkyl group is optionally substituted by one or more halogen groups and in which one or more non-adjacent alkyl -CH₂- groups are independently optionally replaced by a group selected from -O-, -CO-, -CO-O-,
 -O-CO-, -CH=CH- and -C≡C-.

20 It is especially preferred that D is selected from pyrimidine-2,5-diyl, pyridine-2,5-diyl, 2,5-furanylene, 1,4- or 2,6-naphthylene and phenylene, which is optionally substituted by a C₁₋₆ straight-chain or branched alkyl residue, which alkyl group is optionally substituted by one or more fluorine atoms, and wherein one or
 25 more non-adjacent alkyl -CH₂- groups are independently optionally replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO- and -CH=CH-.

By the term "phenylene" it should be understood to include 1,2-, 1,3- or 1,4-phenylene, which is optionally substituted. It is preferred that the phenylene group is either a 1,3- or a 1,4-phenylene. 1,4-phenylene groups are especially preferred.

Preferred groups E are selected from $-OR^3$ and $-NR^4R^5$, wherein R^3 and R^4 represent a C_{1-18} cyclic, straight-chain or branched alkyl residue, which is optionally substituted by one or more halogen atoms, and in which one or more non-adjacent alkyl $-CH_2-$ groups are independently optionally replaced by $-O-$ or $-CH=CH-$, wherein R^5 is selected from a hydrogen atom or a C_{1-18} cyclic, straight-chain or branched alkyl residue, which is optionally substituted by one or more halogen groups and which one or more non-adjacent alkyl $-CH_2-$ groups are optionally independently replaced by $-O-$ or $-CH=CH-$, or R^4 and R^5 together to form a C_{5-8} alicyclic ring.

It is especially preferred that E is selected from the group comprising $-OR^3$ or $-NHR^4$, wherein R^3 and R^4 represent a C_{1-18} cyclic, straight-chain or branched alkyl residue which is optionally substituted by one or more fluorine atoms and in which one or more non-adjacent alkyl $-CH_2-$ groups are independently optionally replaced by $-O-$.

Preferred groups B1 to B4 are groups of formula II where $n^4 + n^5 \leq 1$.

It is preferred that each of the groups C^1 and C^2 comprising the groups B^1 to B^4 are selected from cyclohexane-1,4-diyl, pyrimidine-2,5-diyl, pyridine-2,5-diyl, 1,4- or 2,6-naphthylene and phenylene, which is optionally substituted by one or more groups selected from fluorine, chlorine, cyano and a C_{1-12} cyclic, straight-chain or branched alkyl residue, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkyl $-CH_2-$ groups are optionally independently replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-CH=CH-$, $-C\equiv C-$ and $-O-CO-O-$.

It is especially preferred that the groups C^1 and C^2 are selected from cyclohexane-1,4-diyl, pyrimidine-2,5-diyl, pyridine-2,5-diyl, 2,6-naphthylene and

phenylene, which is optionally substituted by one or more fluorine atoms or a C₁₋₈ straight-chain or branched alkyl residue, which is optionally substituted by one or more fluorine atoms, and in which one or more non-adjacent alkyl -CH₂- groups are independently optionally replaced by a group selected from -O-, -CO-, -CO-O-,
 5 -O-CO- and -CH=CH-.

It is preferred that the groups S¹ to S³ are selected from a single covalent bond, -O-, -CO-O-, -O-CO-, -NR¹-, -NR¹-CO-, -CO-NR¹-, -NR¹-CO-O-,
 10 -O-CO-NR¹-, -NR¹-CO-NR¹-, -CH=CH-, -C≡C-, -O-CO-O- and a straight-chain or branched alkylene group, which is optionally substituted by one or more groups selected from fluorine, chlorine and cyano and in which two or three non-adjacent alkylene -CH₂- group are independently optionally replaced by a group selected from -O-, -CO-O-, -O-CO-, -NR¹-, -NR¹-CO-, -CO-NR¹-,
 15 -NR¹-CO-O-, -O-CO-NR²-, -NR¹-CO-NR¹-, -CH=CH-, -C≡C-, -O-CO-O- and -Si(CH₃)₂-O-Si(CH₃)₂-, wherein R¹ is as defined above and with the proviso that firstly, the total number of chain carbon atoms in the alkylene group does not exceed 24 and secondly, when the repeating monomer unit M is linked to B⁴ via a nitrogen atom or a oxygen atom S¹, S² and S³ are not -O-, -CO-O-, -O-CO-,
 -NR¹-, -NR¹-CO-, -CO-NR¹-, -NR¹-CO-O-, -O-CO-NR¹-,
 -NR¹-CO-NR¹-, -CH=CH-, -C≡C- or -O-CO-O-.

It is more preferred that S¹ to S³ are selected from -CO-O-, -O-CO-,
 20 -(CH₂)_r-, -(CH₂)_r-O-, -(CH₂)_r-CO-, -(CH₂)_r-CO-O-, -(CH₂)_r-O-CO-,
 -(CH₂)_r-CO-NR¹-, -(CH₂)_r-NR¹-CO-, -(CH₂)_r-NR¹-, -O-(CH₂)_r-,
 -CO-O-(CH₂)_r-, -O-CO-(CH₂)_r-, -NR¹-CO-(CH₂)_r-, -CO-NR¹-(CH₂)_r-,
 -NR¹-(CH₂)_r-, -O-(CH₂)_r-CO-O-, -O-(CH₂)_r-O-CO-,
 25 -O-(CH₂)_r-CO-NR¹-, -O-(CH₂)_r-NR¹-, -O-(CH₂)_r-O-, -O-(CH₂)_r-NR¹-CO-,
 -NR¹-(CH₂)_r-CO-O-, -NR¹-(CH₂)_r-O-, -NR¹-(CH₂)_r-NR¹-,

$-\text{NR}^1-(\text{CH}_2)_r-\text{O}-\text{CO}-$, $-\text{CO}-\text{NR}^1-(\text{CH}_2)_r-\text{O}-$, $-\text{CO}-\text{NR}^1-(\text{CH}_2)_r-\text{NR}^1-$,
 $-\text{CO}-\text{NR}^1-(\text{CH}_2)_r-\text{O}-\text{CO}-$, $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{CO}-$, $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{O}-$,
 $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{NR}^2-$, $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{CO}-\text{NR}^1-$,
 $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{NR}^1-\text{CO}-$, $-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-$, $-(\text{CH}_2)_r-\text{CO}-\text{O}-(\text{CH}_2)_s-$,
5 $-(\text{CH}_2)_r-\text{O}-\text{CO}-(\text{CH}_2)_s-$, $-(\text{CH}_2)_r-\text{NR}^1-\text{CO}-(\text{CH}_2)_s-$, $-(\text{CH}_2)_r-\text{NR}^1-$
 $\text{CO}-\text{O}-(\text{CH}_2)_s-$, $-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-(\text{CH}_2)_r-\text{CO}-\text{O}-(\text{CH}_2)_s-\text{O}-$,
 $-(\text{CH}_2)_r-\text{O}-\text{CO}-(\text{CH}_2)_s-\text{O}-$, $-(\text{CH}_2)_r-\text{NR}^1-\text{CO}-(\text{CH}_2)_s-\text{O}-$, $-(\text{CH}_2)_r-\text{NR}^1-$
 $\text{CO}-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-\text{O}-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-$, $-\text{O}-(\text{CH}_2)_r-\text{CO}-\text{O}-(\text{CH}_2)_s-$,
 $-\text{O}-(\text{CH}_2)_r-\text{NR}^1-\text{CO}-(\text{CH}_2)_s-$, $-\text{O}-(\text{CH}_2)_r-\text{NR}^1-\text{CO}-\text{O}-(\text{CH}_2)_s-$, $-\text{O}-(\text{CH}_2)_r-$
10 $\text{COO}-(\text{CH}_2)_s-\text{O}-$, $-\text{O}-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-\text{O}-(\text{CH}_2)_r-\text{NR}^1-\text{CO}-(\text{CH}_2)_s-\text{O}-$,
 $-\text{O}-(\text{CH}_2)_r-\text{NR}^1-\text{CO}-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-\text{CO}-\text{O}-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-$ and $-\text{CO}-\text{O}-$
 $(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-\text{O}-$, wherein R^1 is as defined above, r and s each represent an
integer from 1 to 20, preferably from 2 to 12, and $r + s \leq 21$, preferably ≤ 15 .

By the terms $-(\text{CH}_2)_r-$ and $-(\text{CH}_2)_s-$ it should be understood to include
15 straight-chain or branched alkylene groupings containing r or s carbon atoms
respectively. Optional substituents include alkyl, aryl, cycloalkyl, amino, cyano,
epoxy, halogen, hydroxy, nitro and oxo.

It is especially preferred that S^1 to S^3 are selected from $-(\text{CH}_2)_r-$,
 $-(\text{CH}_2)_r-\text{O}-$, $-(\text{CH}_2)_r-\text{CO}-\text{O}-$, $-(\text{CH}_2)_r-\text{O}-\text{CO}-$, $-(\text{CH}_2)_r-\text{CO}-\text{NH}-$,
20 $-(\text{CH}_2)_r-\text{NH}-\text{CO}-$, $-\text{O}-(\text{CH}_2)_r-$, $-\text{CO}-\text{O}-(\text{CH}_2)_r-$, $-\text{CO}-\text{NH}-(\text{CH}_2)_r-$,
 $-\text{O}-\text{CO}-(\text{CH}_2)_r-$, $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{CO}-\text{O}-$, $-\text{O}-(\text{CH}_2)_r-\text{O}-\text{CO}-$, $-\text{O}(\text{CH}_2)_r-$
 $\text{CO}-\text{NH}-$, $-\text{O}-(\text{CH}_2)_r-\text{NH}-\text{CO}-$, $-\text{CO}-\text{O}-(\text{CH}_2)_r-\text{O}-$, $-\text{CO}-\text{NH}-(\text{CH}_2)_r-\text{O}-$,
 $-\text{O}-(\text{CH}_2)_r-\text{O}-$, $-(\text{CH}_2)_r-\text{NH}-\text{CO}-(\text{CH}_2)_s-$, $-(\text{CH}_2)_r-\text{NH}-\text{CO}-\text{O}-(\text{CH}_2)_s-$,
 $-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-(\text{CH}_2)_r-\text{NH}-\text{CO}-(\text{CH}_2)_s-\text{O}-$, $-(\text{CH}_2)_r-\text{NH}-$
25 $\text{CO}-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-\text{O}-(\text{CH}_2)_r-\text{NH}-\text{CO}-(\text{CH}_2)_s-$, $-\text{O}-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-\text{O}-$,

$-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-\text{CO}-\text{O}-(\text{CH}_2)_r-\text{O}-(\text{CH}_2)_s-\text{O}-$, $-\text{O}-(\text{CH}_2)_r-\text{NH}-\text{CO}-(\text{CH}_2)_s-\text{O}-$ and $-\text{O}-\text{CO}-(\text{CH}_2)_r-\text{NH}-\text{CO}-(\text{CH}_2)_s-\text{O}-$, wherein r and s each represent an integer from 2 to 12 and $r + s \leq 15$.

Examples of preferred the preferred groups S^1 to S^3 include 1,2-ethylene,
 5 1,3-propylene, 1,4-butylene, 1,5-pentylene, 1,6-hexylene, 1,7-heptylene, 1,8-octylene,
 1,9-nonylene, 1,10-decylene, 1,11-undecylene, 1,12-dodecylene, 3-methyl-
 -1,4-butylene, 3-propyleneoxy, 3-propyleneoxycarbonyl, 2-ethylenecarbonyloxy,
 4-butyleneoxy, 4-butyleneoxycarbonyl, 3-propylenecarbonyloxy, 5-pentyleneoxy,
 5-pentyleneoxycarbonyl, 4-butylenecarbonyloxy, 6-hexyleneoxy, 6-hexyleneoxyc-
 10 arbonyl, 5-pentylenecarbonyloxy, 7-heptyleneoxy, 7-heptyleneoxycarbonyl,
 6-hexylenecarbonyloxy, 8-octyleneoxy, 8-octyleneoxycarbonyl, 7-heptylenecarbonyl-
 oxy, 9-nonyleneoxy, 9-nonyleneoxycarbonyl, 8-octylenecarbonyloxy, 10-decylene-
 oxy, 10-decyleneoxycarbonyl, 9-nonylenecarbonyloxy, 11-undecyleneoxy,
 11-undecyleneoxycarbonyl, 10-decylenecarbonyloxy, 12-dodecyleneoxy,
 15 12-dodecyleneoxycarbonyl, 11-undecylenecarbonyloxy, 3-propyleneiminocarbonyl,
 4-butyleneiminocarbonyl, 5-pentyleneiminocarbonyl, 6-hexyleneiminocarbonyl,
 7-heptyleneiminocarbonyl, 8-octyleneiminocarbonyl, 9-nonyleneiminocarbonyl,
 10-decyleneiminocarbonyl, 11-undecyleneiminocarbonyl, 12-dodecyleneimino-
 carbonyl, 2-ethylenecarbonylimino, 3-propylenecarbonylimino, 4-butylenecarbonyl-
 20 imino, 5-pentylenecarbonylimino, 6-hexylenecarbonylimino, 7-heptylenecarbonyl-
 imino, 8-octylenecarbonylimino, 9-nonylenecarbonylimino, 10-decylenecarbonyl-
 imino, 11-undecylenecarbonylimino, 6-(3-propyleneiminocarbonyloxy)hexylene,
 6-(3-propyleneoxy)hexylene, 6-(3-propyleneoxy)hexyleneoxy, 6-(3-propyleneimino-
 carbonyloxy)hexyleneoxy, 6-(3-propyleneiminocarbonyl)hexyl, 6-(3-propyleneimino-
 25 carbonyl)hexyloxy, 1,2-ethylenedioxy, 1,3-propylenedioxy, 1,4-butylenedioxy,
 1,5-pentylenedioxy, 1,6-hexylenedioxy, 1,7-heptylenedioxy, 1,8-octylenedioxy,
 1,9-nonylenedioxy, 1,10-decylenedioxy, 1,11-undecylenedioxy, 1,12-dodecylene-
 dioxy and the like.

It is preferred that the unit of formula (I) comprises at least 50% of the monomer building blocks, which form the main chain of a photoactive polymer. It is especially preferred that the unit of formula (I) comprises at least 70% of the monomer building blocks forming the photoactive polymer.

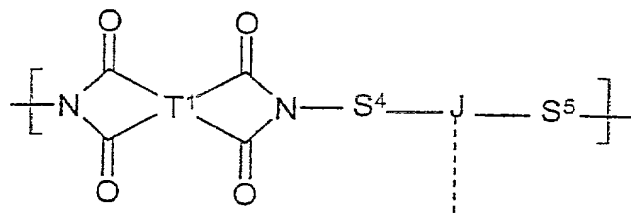
5 The repeating monomer unit M represents part of a homopolymer or a copolymer. It is preferred that M forms part of a co-polymer. By the term "copolymer" it is to be understood to include statistical copolymers.

10 The repeating monomer units M are preferably selected from acrylate; methacrylate; 2-chloroacrylate; 2-phenylacrylate; acrylamide, methacrylamide, 2-chloroacrylamide and 2-phenylacrylamide, the nitrogen atom of which is optionally substituted by a lower alkyl group; vinyl ether; vinyl ester; a styrene derivative; siloxane; imide; amic acid; amic acid esters; amidimide; maleic acid derivatives and fumaric acid derivatives.

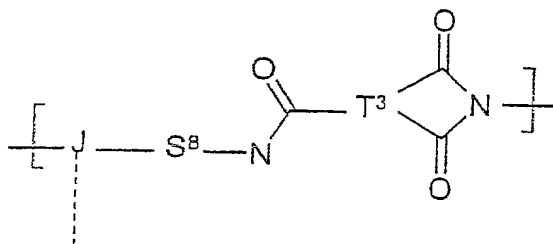
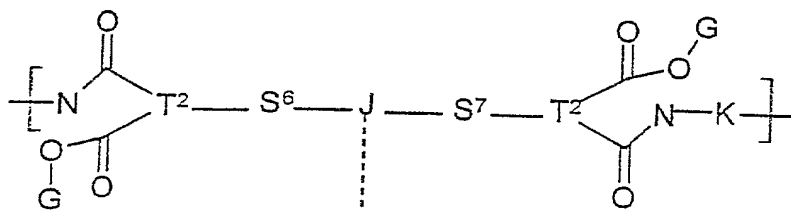
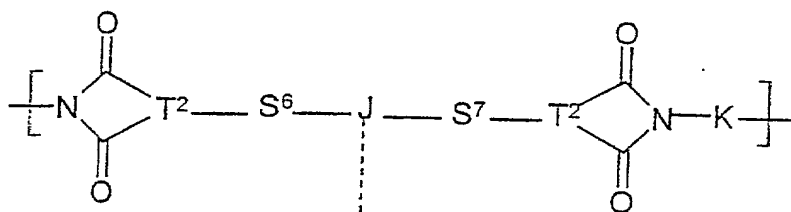
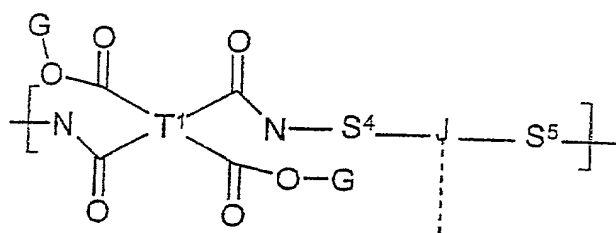
15 It is more preferred that the repeating monomer unit M is selected from acrylate; methacrylate; acrylamide and methacrylamide the nitrogen atom of which is optionally substituted by a lower alkyl group; vinyl ether; vinyl ester; a styrene derivative, imide, amic acid, amic acid esters and amidimide.

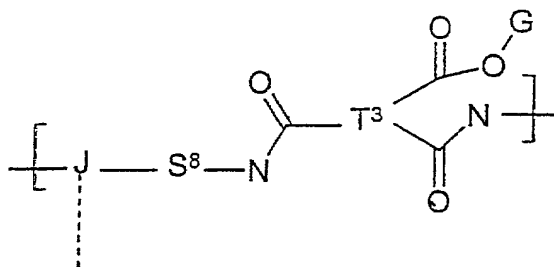
20 It is especially preferred that the repeating monomer unit M is selected from acrylate, methacrylate, a styrene derivative, imide, amic acid, amic acid ester, and amidimide.

When the monomer unit M is an imide group, it is preferably selected from structures of the general formulae VI, VII, VIII, IX, X and XI:



VI





XI

wherein the broken line symbolises the linkage to B⁴

T¹ represents a tetravalent organic radical;

T², T³ each independently represent a trivalent aromatic or alicyclic group which is optionally substituted by a group selected from fluorine, chlorine, cyano and a C₁₋₁₈ cyclic, straight-chain or branched alkyl residue, which is optionally substituted by one or more halogen groups and in which one or more non-adjacent alkyl -CH₂- groups are independently optionally replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -CH=CH- and -C≡C-,

S⁴ to S⁸ are each independently selected from a single covalent bond and a C₁₋₂₄ straight-chain or branched alkylene residue, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkylene -CH₂- groups are, independently, optionally be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -Si(CH₃)₂-O-Si(CH₃)₂-, -NR¹-, -NR¹-CO-, -CO-NR¹-, -NR¹-CO-O-, -O-CO-NR¹-,

$-\text{NR}^1-\text{CO}-\text{NR}^1-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$ and $-\text{O}-\text{CO}-\text{O}-$, wherein R^1 is as defined above;

J is selected from the group comprising a nitrogen atom, a group $-\text{CR}^1-$ and an aromatic or alicyclic divalent, trivalent or tetravalent group, which is optionally substituted by one or more groups selected from fluoro, chloro, cyano and a C_{1-18} cyclic, straight-chain or branched alkyl residue which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent $-\text{CH}_2-$ groups are, independently, optionally, replaced by a group selected from $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-$, $-\text{CH}=\text{CH}-$ and $-\text{C}\equiv\text{C}-$, wherein R^1 is as defined above;

K represents an aliphatic, alicyclic or aromatic divalent radical; and

G represents a hydrogen atom or a monovalent organic group.

By the term "aliphatic" it should be understood to include saturated and unsaturated, straight-chain and branched alkyl groups, which may be optionally substituted and in which one or more non-adjacent $-\text{CH}_2-$ groups are replaced by one or more heteroatoms. Optional substituents include alkyl, aryl, cycloalkyl, amino, cyano, epoxy, halogen, hydroxy, nitro and oxo. Examples of heteroatoms that can replace the one or more $-\text{CH}_2-$ groups include nitrogen, oxygen and sulfur. Replacement nitrogen atoms may be further substituted with groups such as alkyl, aryl and cycloalkyl.

The tetravalent organic radical T^1 is preferably derived from an aliphatic, alicyclic or aromatic tetracarboxylic acid dianhydride. Alicyclic or aliphatic tetracarboxylic acid anhydrides are preferably selected from butanetetracarboxylic acid dianhydride, ethylenemaleic acid dianhydride, 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride, 1,2,3,4-cyclopentanetetracarboxylic acid dianhydride,

2,3,5-tricarboxycyclopentylacetic acid dianhydride, 3,5,6-tricarboxynorbornylacetic acid dianhydride, 2,3,4,5-tetrahydrofuran-tetracarboxylic acid dianhydride, 4-(2,5-dioxotetrahydrofuran-3-yl)tetrahydronaphthalene-1,2-dicarboxylic acid dianhydride, 5-(2,5-dioxotetrahydrofuran-3-yl)-3-methyl-3-cyclohexene-1,2-dicarboxylic acid dianhydride, bicyclo[2.2.2]oct-7-ene-2,3,5,6-tetracarboxylic acid dianhydride, bicyclo[2.2.2]octane-2,3,5,6-tetracarboxylic acid dianhydride and 1,8-dimethylbicyclo[2.2.2]oct-7-ene-2,3,5,6-tetracarboxylic acid dianhydride.

Aromatic tetracarboxylic acid dianhydrides are preferably selected from pyromellitic acid dianhydride, 3,3',4,4'-benzophenonetetracarboxylic acid dianhydride, 4,4'-oxydiphthalic acid dianhydride, 3,3',4,4'-diphenylsulfonetetracarboxylic acid dianhydride, 1,4,5,8-naphthalenetetracarboxylic acid dianhydride, 2,3,6,7-naphthalenetetracarboxylic acid dianhydride, 3,3',4,4'-dimethyldiphenylsilane-tetracarboxylic acid dianhydride, 3,3',4,4'-tetraphenylsilane-tetracarboxylic acid dianhydride, 1,2,3,4-furantetracarboxylic acid dianhydride, 4,4'-bis(3,4-dicarboxyphenoxy)diphenyl sulfide dianhydride, 4,4'-bis(3,4-dicarboxyphenoxy)diphenyl sulfone dianhydride, 4,4'-bis(3,4-dicarboxyphenoxy)diphenylpropane dianhydride, 3,3',4,4'-biphenyltetracarboxylic acid dianhydride, ethylene glycol bis(trimellitic acid) dianhydride, 4,4'-(1,4-phenylene)bis(phthalic acid) dianhydride, 4,4'-(1,3-phenylene)bis(phthalic acid) dianhydride, 4,4'-(hexafluoroisopropylidene)diphthalic acid dianhydride, 4,4'-oxydi(1,4-phenylene)bis(phthalic acid) dianhydride and 4,4'-methylenedi(1,4-phenylene)bis(phthalic acid) dianhydride.

It is especially preferred that the tetracarboxylic acid dianhydrides used to form the tetravalent organic radical T^1 are selected from 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride, 1,2,3,4-cyclopentanetetracarboxylic acid dianhydride, 2,3,5-tricarboxycyclopentylacetic acid dianhydride, 5-(2,5-dioxotetrahydrofuran-3-yl)-3-methyl-3-cyclohexene-1,2-dicarboxylic acid dianhydride, 4-(2,5-dioxotetrahydrofuran-3-yl)tetrahydronaphthalene-1,2-dicarboxylic acid dianhydride, 4,4'-(hexafluoroisopropylidene)diphthalic acid dianhydride and bicyclo[2.2.2]oct-7-ene-2,3,5,6-tetracarboxylic acid dianhydride.

Each of the groups T^2 and T^3 can be derived from an aliphatic, alicyclic or aromatic dicarboxylic acid anhydride.

The groups T^2 and T^3 are preferably trivalent aromatic or alicyclic groups, the three valencies of which are distributed between three different carbon atoms, with the proviso that two of the valencies are located at adjacent carbon atoms. It is especially preferred that the groups T^2 and T^3 are trivalent benzene derivatives.

The group S^4 is preferably selected from a single covalent bond, $-(CH_2)_r-$, $-(CH_2)_r-O-$, $-(CH_2)_r-CO-$, $-(CH_2)_r-CO-O-$, $-(CH_2)_r-O-CO-$, $-(CH_2)_r-CO-NR^1-$, $-(CH_2)_r-NR^1-CO-$, $-(CH_2)_r-NR^1-$, $-(CH_2)_r-O-(CH_2)_s-$, $-(CH_2)_r-CO-O-(CH_2)_s-$, $-(CH_2)_r-O-CO-(CH_2)_s-$, $-(CH_2)_r-NR^1-CO-(CH_2)_s-$, $-(CH_2)_r-NR^1-CO-O-(CH_2)_s-$, $-(CH_2)_r-O-(CH_2)_s-O-$, $-(CH_2)_r-CO-O-(CH_2)_s-O-$, $-(CH_2)_r-O-CO-(CH_2)_s-O-$, $-(CH_2)_r-NR^1-CO-(CH_2)_s-O-$, $-(CH_2)_r-NR^1-CO-O-(CH_2)_s-O-$, $-(CH_2)_r-O-(CH_2)_s-CO-O-$ and $-(CH_2)_r-O-(CH_2)_s-O-CO-$, wherein R^1 is as defined herein above; r and s each represent an integer from 1 to 20; and $r + s \leq 21$. It is more preferred that r and s each represent an integer from 2 to 12. It is especially preferred that $r + s \leq 15$.

Examples of preferred groups S^4 include 1,2-ethylene, 1,3-propylene, 1,4-butylene, 1,5-pentylene, 1,6-hexylene, 1,7-heptylene, 1,8-octylene, 1,9-nonylene, 1,10-decylene, 1,11-undecylene, 1,12-dodecylene, 3-methyl-1,4-butylene, 3-propyleneoxy, 3-propyleneoxycarbonyl, 2-ethylenecarbonyloxy, 4-butyleneoxy, 4-butyleneoxycarbonyl, 3-propylenecarbonyloxy, 5-pentyleneoxy, 5-pentyleneoxycarbonyl, 4-butylenecarbonyloxy, 6-hexyleneoxy, 6-hexyleneoxycarbonyl, 5-pentylenecarbonyloxy, 7-heptyleneoxy, 7-heptyleneoxycarbonyl, 6-hexylenecarbonyloxy, 8-octyleneoxy, 8-octyleneoxycarbonyl, 7-heptylenecarbonyloxy, 9-nonyleneoxy, 9-nonyleneoxycarbonyl, 8-octylenecarbonyloxy, 10-decyleneoxy, 10-decyleneoxycarbonyl, 9-nonylenecarbonyloxy, 11-undecyleneoxy, 11-undecylene-

oxycarbonyl, 10-decylenecarbonyloxy, 12-dodecyleneoxy, 12-dodecylene-
 oxycarbonyl, 11-undecylenecarbonyloxy, 3-propyleneiminocarbonyl, 4-butylene-
 iminocarbonyl, 5-pentyleneiminocarbonyl, 6-hexyleneiminocarbonyl, 7-heptylene-
 iminocarbonyl, 8-octyleneiminocarbonyl, 9-nonyleneiminocarbonyl, 10-decylene-
 5 iminocarbonyl, 11-undecyleneiminocarbonyl, 12-dodecyleneiminocarbonyl,
 2-ethylenecarbonylimino, 3-propylenecarbonylimino, 4-butylenecarbonylimino,
 5-pentylenecarbonylimino, 6-hexylenecarbonylimino, 7-heptylenecarbonylimino,
 8-octylenecarbonylimino, 9-nonylenecarbonylimino, 10-decylenecarbonylimino,
 11-undecylenecarbonylimino, 6-(3-propyleneiminocarbonyloxy)hexylene,
 10 6-(3-propyleneoxy)hexylene, 6-(3-propyleneoxy)hexyleneoxy, 6-(3-propyleneimino-
 carbonyloxy)hexyleneoxy, 6-(3-propyleneiminocarbonyl)hexylene, 6-(3-propylene-
 iminocarbonyl)hexyleneoxy and the like.

The groups S^5 and S^8 are preferably selected from a single bond, $-(CH_2)_r-$,
 $-O-(CH_2)_r-$, $-CO-(CH_2)_r-$, $-CO-O-(CH_2)_r-$, $-O-CO-(CH_2)_r-$,
 15 $-NR^1-CO-(CH_2)_r-$, $-NR^1-(CH_2)_r-$, $-CO-NR^1-(CH_2)_r-$, $-NR^1-CO-(CH_2)_r-$,
 $-O-(CH_2)_r-O-(CH_2)_s-$, $-(CH_2)_r-CO-O-(CH_2)_s-$, $-(CH_2)_r-O-CO-(CH_2)_s-$,
 $-(CH_2)_r-NR^1-CO-(CH_2)_s-$, $-(CH_2)_r-NR^1CO-O-(CH_2)_s-$,
 $-O-(CH_2)_r-O-(CH_2)_s-$, $-O-(CH_2)_rCO-O-(CH_2)_s-$,
 $-O-(CH_2)_r-O-CO-(CH_2)_s-$, $-O-(CH_2)_r-NR^1-CO-(CH_2)_s-$, $-O-(CH_2)_r-NR^1-$
 20 $CO-O-(CH_2)_s-$, $-O-CO-(CH_2)_r-O-(CH_2)_s-$ and $-CO-O-(CH_2)_r-O-(CH_2)_s-$,
 wherein R^1 is defined as herein above; r and s each represent an integer from 1 to 20;
 and $r + s \leq 21$. It is more preferred that r and s each represent an integer from 2 to 12.
 It is further preferred that $r + s \leq 15$.

Examples of preferred groups S^5 and S^8 include 1,2-ethylene, 1,3-propylene,
 25 1,4-butylene, 1,5-pentylene, 1,6-hexylene, 1,7-heptylene, 1,8-octylene, 1,9-nonylene,
 1,10-decylene, 1,11-undecylene, 1,12-dodecylene, 3-methyl-1,4-butylene, 2-oxy-
 ethylene, 3-oxypropylene, 4-oxybutylene, 5-oxypropylene, 6-oxyhexylene, 7-oxy-

heptylene, 8-oxyoctylene, 9-oxynonylene, 10-oxydecylene, 11-oxyundecylene, 12-oxydodecylene, 2-(oxycarbonyl)ethylene, 3-(oxycarbonyl)propylene, 4-(oxycarbonyl)butylene, 5-(oxycarbonyl)pentylene, 6-(oxycarbonyl)hexylene, 7-(oxycarbonyl)heptylene, 8-(oxycarbonyl)octylene, 9-(oxycarbonyl)nonylene, 10-(oxycarbonyl)decylene, 11-(oxycarbonyl)undecylene, 12-(oxycarbonyl)-
 5 dodecylene, 2-(carbonyloxy)ethylene, 3-(carbonyloxy)propylene, 4-(carbonyloxy)-butylene, 5-(carbonyloxy)pentylene, 6-(carbonyloxy)hexylene, 7-(carbonyloxy)-heptylene, 8-(carbonyloxy)octylene, 9-(carbonyloxy)nonylene, 10-(carbonyloxy)-decylene, 11-(carbonyloxy)undecylene, 12-(carbonyloxy)dodecylene, 2-(carbonyl-
 10 imino)ethylene, 3-(carbonylimino)propylene, 4-(carbonylimino)butylene, 5-(carbonylimino)pentylene, 6-(carbonylimino)hexylene, 7-(carbonylimino)heptylene, 8-(carbonylimino)octylene, 9-(carbonylimino)nonylene, 10-(carbonylimino)decylene, 11-(carbonylimino)undecylene, 12-(carbonylimino)dodecylene, 2-iminoethylene, 3-iminopropylene, 4-iminobutylene, 5-iminopentylene, 6-iminohexylene, 7-imino-
 15 heptylene, 8-iminooctylene, 9-iminononylene, 10-iminodecylene, 11-imino-undecylene, 12-iminododecylene, 2-iminocarbonylethylene, 3-imino-carbonylpropylene, 4-iminocarbonylbutylene, 5-iminocarbonylpentylene, 6-imino-carbonylhexylene, 7-iminocarbonylheptylene, 8-iminocarbonyloctylene, 9-imino-carbonylnonylene, 10-iminocarbonyldecylene, 11-iminocarbonylundecylene, 12-iminocarbonyldodecylene, 2-(2-ethyleneoxy)ethylene, 2-(3-propyleneoxy)-
 20 ethylene, 6-(4-butyleneoxy)hexylene, 2-(2-ethyleneiminocarbonyl)ethylene, 2-(3-propyleneiminocarbonyl)ethylene, 6-(4-butyleneiminocarbonyl)hexylene, 6-(3-propyleneiminocarbonyloxy)hexylene, 6-(3-propyleneiminocarbonyl)hexylene and the like.

25 The groups S^6 and S^7 are preferably selected from a single bond, $-(CH_2)_I-$, $-(CH_2)_I-O-$, $-(CH_2)_I-CO-$, $-(CH_2)_I-CO-O-$, $-(CH_2)_I-O-CO-$, $-(CH_2)_I-CO-NR^1-$, $-(CH_2)_I-NR^1-CO-$, $-(CH_2)_I-NR^1-$, $-O-(CH_2)_I-$, $-CO-O-(CH_2)_I-$, $-O-CO-(CH_2)_I-$, $-NR^1-CO-(CH_2)_I-$, $-CO-NR^1-(CH_2)_I-$, $-NR^1-(CH_2)_I-$,

$-O-(CH_2)_r-CO-O-$, $-O-(CH_2)_r-O-CO-$, $-O-(CH_2)_r-CO-NR^1-$, $-O-(CH_2)_r-NR^1-$,
 $-O-(CH_2)_r-O-$, $-O-(CH_2)_r-NR^1-CO-$, $-NR^1-(CH_2)_r-CO-O-$, $-NR^1-(CH_2)_r-O-$,
 $-NR^1-(CH_2)_r-NR^1-$, $-NR^1-(CH_2)_r-O-CO-$, $-CO-NR^1(CH_2)_r-O-$,
 $-CO-NR^1-(CH_2)_r-NR^1-$, $-CO-NR^1-(CH_2)_r-O-CO-$, $-O-CO(CH_2)_r-CO-$,
5 $-O-CO-(CH_2)_r-O-$, $-O-CO-(CH_2)_r-NR^1-$, $-O-CO-(CH_2)_r-CO-O-$,
 $-O-CO-(CH_2)_r-CO-NR^1-$, $-O-CO-(CH_2)_r-NR^1-CO-$, $-(CH_2)_r-O-(CH_2)_s-$,
 $-(CH_2)_r-CO-O-(CH_2)_s-$, $-(CH_2)_r-O-CO-(CH_2)_s-$, $-(CH_2)_r-NR^1-CO-(CH_2)_s-$,
 $-(CH_2)_r-NR^1-CO-O-(CH_2)_s-$, $-(CH_2)_r-O-(CH_2)_s-O-$, $-(CH_2)_r-CO-O-(CH_2)_s-O-$,
 $-(CH_2)_r-O-CO-(CH_2)_s-O-$, $-(CH_2)_r-NR^1-CO-(CH_2)_s-O-$, $-(CH_2)_r-NR^1-$
10 $CO-O-(CH_2)_s-O-$, $-O-(CH_2)_r-O-(CH_2)_s-$, $-O-(CH_2)_r-CO-O-(CH_2)_s-$,
 $-O-(CH_2)_r-NR^1-CO-(CH_2)_s-$, $-O-(CH_2)_r-NR^1-CO-O-(CH_2)_s-$, $-O-(CH_2)_r-CO-$
 $O-(CH_2)_s-O-$, $-O-(CH_2)_r-O-(CH_2)_s-O-$, $-O-(CH_2)_r-NR^1-CO-(CH_2)_s-O-$,
 $-O-(CH_2)_r-NR^1-CO-O-(CH_2)_s-O-$, $-CO-O-(CH_2)_r-O-(CH_2)_s-$, $-CO-O-(CH_2)_r-$
 $O-(CH_2)_s-O-$, wherein R^1 is defined as herein above; r and s each represent an integer
15 from 1 to 20; and $r + s \leq 21$. It is more preferred that r and s each represent an integer
from 2 to 12. It is especially preferred that $r + s \leq 15$.

Examples of preferred groups S^6 and S^7 include 1,2-ethylene, 1,3-propylene,
 1,4-butylene, 1,5-pentylene, 1,6-hexylene, 1,7-heptylene, 1,8-octylene, 1,9-nonylene,
 1,10-decylene, 1,11-undecylene, 1,12-dodecylene, 3-methyl-1,4-butylene,
 20 3-propyleneoxy, 3-propyleneoxycarbonyl, 2-ethylenecarbonyloxy, 4-butyleneoxy,
 4-butyleneoxycarbonyl, 3-propylenecarbonyloxy, 5-pentyleneoxy, 5-pentylene-
 oxycarbonyl, 4-butylenecarbonyloxy, 6-hexyleneoxy, 6-hexyleneoxycarbonyl,
 5-pentylenecarbonyloxy, 7-heptyleneoxy, 7-heptyleneoxycarbonyl, 6-hexylene-
 carbonyloxy, 8-octyleneoxy, 8-octyleneoxycarbonyl, 7-heptylenecarbonyloxy,
 25 9-nonyleneoxy, 9-nonyleneoxycarbonyl, 8-octylenecarbonyloxy, 10-decyleneoxy,

10-decylenecarbonyl, 9-nonylenecarbonyloxy, 11-undecylenecarbonyl, 10-decylenecarbonyloxy, 12-dodecylenecarbonyl, 11-undecylenecarbonyloxy, 3-propyleneiminocarbonyl, 4-butyleneiminocarbonyl, 5-pentyleneiminocarbonyl, 6-hexyleneiminocarbonyl, 7-heptyleneiminocarbonyl, 8-octyleneiminocarbonyl, 9-nonyleneiminocarbonyl, 10-decylenecarbonyl, 11-undecylenecarbonyl, 12-dodecylenecarbonyl, 2-ethylenecarbonylimino, 3-propylenecarbonylimino, 4-butylenecarbonylimino, 5-pentylene-carbonylimino, 6-hexylenecarbonylimino, 7-heptylenecarbonylimino, 8-octylene-carbonylimino, 9-nonylenecarbonylimino, 10-decylenecarbonylimino, 11-undecylenecarbonylimino, 6-(3-propyleneiminocarbonyloxy)hexylene, 6-(3-propyleneoxy)-hexylene, 6-(3-propyleneoxy)hexyleneoxy, 6-(3-propyleneiminocarbonyloxy)-hexyleneoxy, 6-(3-propyleneiminocarbonyl)hexyl, 6-(3-propyleneiminocarbonyl)-hexyloxy, 1,2-ethylenedioxy, 1,3-propylenedioxy, 1,4-butylenedioxy, 1,5-pentylene-dioxy, 1,6-hexylenedioxy, 1,7-heptylenedioxy, 1,8-oxylenedioxy, 1,9-nonylenedioxy, 1,10-decylenedioxy, 1,11-undecylenedioxy, 1,12-dodecylenedioxy and the like.

The aliphatic, alicyclic or aromatic divalent radical K is derivable from aliphatic, alicyclic or aromatic diamines by formal removal of the amino groups. Examples of aliphatic or alicyclic diamines from which the radical K can be derived include ethylenediamine, 1,3-propylenediamine, 1,4-butylenediamine, 1,5-pentylene-diamine, 1,6-hexylenediamine, 1,7-heptylenediamine, 1,8-oxylenediamine, 1,9-nonylenediamine, 1,10-decylenediamine, 1,11-undecylenediamine, 1,12-dodecylenediamine, α,α' -diamino-*m*-xylene, α,α' -diamino-*p*-xylene, (5-amino-2,2,4-trimethylcyclopentyl)methylamine, 1,2-diaminocyclohexane, 4,4'-diamino-dicyclohexylmethane, 1,3-bis(methylamino)cyclohexane, and 4,9-dioxadodecane-1,12-diamine.

Examples of aromatic diamines from which the radical K can be derived include 3,5-diaminobenzoic acid methyl ester, 3,5-diaminobenzoic acid hexyl ester, 3,5-diaminobenzoic acid dodecyl ester, 3,5-diaminobenzoic acid isopropyl ester, 4,4'-methylenedianiline, 4,4'-ethylenedianiline, 4,4'-diamino-3,3'-dimethyldiphenyl-

methane, 3,3',5,5'-tetramethylbenzidine, 4,4'-diaminodiphenyl sulfone, 4,4'-diamino-
 diphenyl ether, 1,5-diaminonaphthalene, 3,3'-dimethyl-4,4'-diaminobiphenyl, 3,4'-diaminodiphenyl ether, 3,3'-diaminobenzophenone, 4,4'-diaminobenzophenone,
 4,4'-diamino-2,2'-dimethylbibenzyl, 2,2-bis[4-(4-aminophenoxy)phenyl] sulfone,
 5 1,4-bis(4-aminophenoxy)benzene, 1,3-bis(4-aminophenoxy)benzene, 1,3-bis(3-amino-
 phenoxy)benzene, 2,7-diaminofluorene, 9,9-bis(4-aminophenyl)fluorene,
 4,4'-methylenebis(2-chloroaniline), 4,4-bis(4-aminophenoxy)biphenyl, 2,2',5,5'-tetra-
 chloro-4,4'-diaminobiphenyl, 2,2'-dichloro-4,4'-diamino-5,5'-dimethoxybiphenyl,
 3,3'-dimethoxy-4,4'-diaminobiphenyl, 4,4'-(1,4-phenyleneisopropylidene)bisani-
 line, 4,4'-(1,3-phenyleneisopropylidene)bisani-
 10 line, 2,2-bis[4-(4-aminophenoxy)phenyl]-
 propane, 2,2-bis[3-(4-aminophenoxy)phenyl]hexafluoropropane, 2,2-bis[3-amino-
 -4-methylphenyl]hexafluoropropane, 2,2-bis(4-aminophenyl)hexafluoropropane,
 2,2'-bis[4-(4-amino-2-trifluoromethylphenoxy)phenyl]hexafluoropropane,
 4,4'-diamino-2,2'-bis(trifluoromethyl)biphenyl, and 4,4'-bis[(4-amino-2-trifluoro-
 15 methyl)phenoxy]-2,3,5,6,2',3',5',6'-octafluorobiphenyl.

The group J may be divalent, trivalent or tetravalent. When J is divalent, it serves to link the groups S⁴ and S⁵, S⁶ and S⁷ and S⁸ and N respectively of the groups VI to XI. It will be appreciated that when J is a divalent group, the monomer unit of which it forms a part is not linked to a side chain group B⁴. When J is a trivalent or a
 20 tetravalent group, it serves to link the monomer unit M, of which it forms a part, to one or two side chain groups B⁴ respectively. It is preferred that the photoactive polymer comprises less than 75 %, of monomer units including a divalent group J, preferably less than 50 % and especially less than 30 %. Monomer units M comprising a trivalent group J are preferred.

25 The building blocks of the formulae VII, IX and XI are amic acid groupings or amic acid ester groupings (i.e. carboxamide-carboxylic acid groupings or carboxamide-carboxylic acid ester groupings) which on the one hand may occur as a result of incomplete imidisation in the polyimide chain. On the other hand, polymers that consist only of building blocks of formulae VII, IX or XI, that is to say polyamic

acids or polyamic acid esters, are important precursors for the preparation of the polyimides according to the invention and are also included in the present invention. Of those polymers which contain groups of formulae VII, IX or XI, preference is given to those in which G is hydrogen, that is to say those which consist exclusively of, or contain some, polyamic acid groups.

The polymers of the invention may be prepared using methods that are known to a person skilled in the art and a second aspect of the invention provides a method of preparing a compound of formula (I) as defined above.

The polymers of formula I, with acrylate, methacrylate and styrene derivative as repeating monomer unit, can be prepared in principle according to two different processes. In addition to the direct polymerisation of pre-finished monomers there exists the possibility of polymer-analogous reaction of reactive photoactive derivatives with functional polymers.

For the direct polymerisation, the monomers and the comonomers are firstly prepared separately from the individual components. The formation of the polymers is subsequently effected in a manner known *per se* under the influence of UV radiation or heat or by the action of radical or ionic catalysts. Potassium peroxodisulfate, dibenzoyl peroxide, azobisisobutyronitrile or di-*tert*-butyl peroxide are examples of radical initiators. Ionic catalysts are alkali-organic compounds such as phenyllithium or naphthylsodium or Lewis acids such as BF_3 , AlCl_3 , SnCl_3 or TiCl_4 . The monomers can be polymerised in solution, suspension, emulsion or substance.

In the second process a polymer of formula I can also be produced in a polymer-analogous reaction from a pre-finished functional polymer and a suitable functionalised photoactive derivative. Many known processes such as, for example, esterification, trans-esterification, amidation or the etherification are suitable for polymer-analogous reactions.

Acrylate, methacrylate and styrene polymers typically have a molecular weight M_w of from 1 000 to 5 000 000, preferably from 5 000 to 2 000 000, and especially from 10 000 to 1 000 000.

Polyamic acids and polyimides of the present invention may be prepared in accordance with known methods, such as those described in *Plast. Eng.* 36 (1996) (Polyimides, fundamentals and applications).

For example, the polycondensation reaction for the preparation of the polyamic acids is carried out in solution in a polar aprotic organic solvent, such as γ -butyrolactone, *N,N*-dimethylacetamide, *N*-methylpyrrolidone or *N,N*-dimethylformamide. In most cases equimolar amounts of the dianhydride and the diamine are used, that is to say one amino group per anhydride group. If it is desired to stabilise the molecular weight of the polymer, it is possible for that purpose to add an excess or a less-than-stoichiometric amount of one of the two components or to add a monofunctional compound in the form of a dicarboxylic acid monoanhydride or in the form of a monoamine. Examples of such monofunctional compounds are maleic acid anhydride, phthalic acid anhydride, aniline and so on. The reaction is carried out preferably at a temperature of less than 100 °C.

The cyclisation of the polyamic acids to form the polyimides can be carried out by heating, that is to say by condensation with removal of water or by other imidisation reactions with reagents. When carried out purely thermally, the imidisation of the polyamic acids is not always complete, that is to say the resulting polyimides may still contain proportions of polyamic acid. The imidisation reactions are generally carried out at a temperature of from 60 to 250 °C, but preferably at less than 200 °C. In order to achieve imidisation at rather lower temperatures there are additionally mixed into the reaction mixture reagents that facilitate the removal of water. Such reagents are, for example, mixtures consisting of acid anhydrides, such as acetic acid anhydride, propionic acid anhydride, phthalic acid anhydride, trifluoroacetic acid anhydride, and tertiary amines, such as triethylamine, trimethylamine, tributylamine, pyridine, *N,N*-dimethylaniline, lutidine, collidine etc.. The amount of reagents used in that case is preferably at least two equivalents of amine and four equivalents of acid anhydride per equivalent of polyamic acid to be condensed.

The imidisation reaction can be carried out before or alternatively only after application to a support. The latter variant is preferred especially when the polyimide in question has poor solubility in the customary solvents.

5 The polyamic acids and the polyimides of the present invention have an intrinsic viscosity preferably in range of 0.05 to 10 dL/g, more preferably 0.05 to 5 dL/g. Herein, the intrinsic viscosity ($\eta_{inh} = \ln \eta_{rel}/C$) is determined by measuring a solution containing a polymer in a concentration of 0.5 g/100 ml for its viscosity at 30 °C using *N*-methyl-2-pyrrolidone as solvent.

10 The polyamic acid chains or polyimide chains of the present invention preferably contain from 2 to 2000 monomer units, especially from 3 to 200.

Additives such as silane-containing compounds and epoxy-containing crosslinking agents may be added to the polymers of the invention in order to improve the adhesion of the polymer to a substrate. Suitable silane-containing compounds are described in *Plast. Eng.* 36 (1996) (Polyimides, fundamentals and applications).
15 Suitable epoxy-containing crosslinking agents include 4,4'-methylenebis-(*N,N*-diglycidylaniline), trimethylolpropane triglycidyl ether, benzene-1,2,4,5-tetracarboxylic acid 1,2:4,5-*N,N'*-diglycidyl diimide, polyethylene glycol diglycidyl ether, *N,N*-diglycidylcyclohexylamine and the like.

Further additives such as photosensitiser, a photoradical generator and/or a cationic photoinitiator may also be added to the polymers of the invention. Suitable photoactive additives include 2,2-dimethoxyphenylethanone, a mixture of diphenylmethanone and *N,N*-dimethylbenzenamine or ethyl-4-(dimethylamino)-benzoate, xanthone, thioxanthone, Irgacure™ 184, 369, 500, 651 and 907 (Ciba),
20 Michler's ketone, triaryl sulfonium salt and the like.

25 The polymers according to the invention may be used alone or in combination with other polymers, oligomers, monomers, photoactive polymers, photoactive oligomers and/or photoactive monomers, depending up on the application to which the polymer layer is to be put. It will therefore be appreciated that by varying the composition of the polymer layer it is possible to control properties such as an

induced pretilt angle, good surface wetting, high voltage holding ratio, a specific anchoring energy etc.

Polymer layers may be readily prepared from the polymers of the present invention and a third aspect of the invention provides a polymer layer comprising a polymer according to the present invention in a cross-linked form. The polymer layer is preferably prepared by applying one or more polymers according to the invention to a support and, after any imidisation step which may be necessary, crosslinking the polymer or polymer mixture by irradiation with linearly polarised light. It is possible to vary the direction of orientation and the tilt angle within the polymer layer by controlling the direction of irradiation of the linearly polarised light. It will be appreciated that by selectively irradiating specific regions of the polymer layer it is possible to align very specific regions of the layer and provide layers with a defined angle of tilt. This orientation and tilt is retained in the polymer layer by the process of cross-linking.

It will be appreciated that the polymer layers of the present invention can also be used as orientation layers for liquid crystals and a preferred embodiment of the third aspect of the invention provides an orientation layer comprising one or more polymers according to the invention in a cross-linked form. Such orientation layers can be used in the manufacture of optical constructional elements, preferably in the production of hybrid layer elements.

The orientation layers are suitably prepared from a solution of the polymer material. The polymer solution is applied to a support optionally coated with an electrode (for example a glass plate coated with indium-tin oxide (ITO)) by a spin coating process, to produce homogeneous layers of 0.05 to 50 μm thickness. The resulting layer is imidised, if required, and may then be selectively orientated by irradiation with a high-pressure mercury vapour lamp, a xenon lamp or a pulsed UV laser, using a polariser and optionally a mask for creating images of structures. The irradiation time is dependent upon the output of the individual lamps and can vary from a few seconds to several hours. The cross-linking reaction can also be carried

out by irradiation using filters that, for example, only allow the radiation suitable for the cross-linking reaction to pass through.

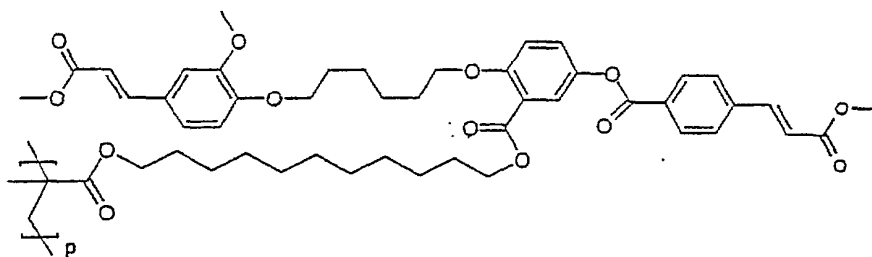
It will be appreciated that the polymer layers of the invention may be used in the production of optical or electro-optical devices having at least one orientation layer as well as unstructured and structured optical elements and multi-layer systems.

A further embodiment of the third aspect of the invention provides an optical or electro-optical device comprising one or more polymers according to the first aspect of the invention in cross-linked form. The electro-optical devices may comprise more than one layers. The or each of the orientation layers may contain one or more regions of different spatial orientation.

The invention will now be described with reference to the following non-limiting examples in which Tg represents the glass temperature, C represents the crystalline phase, N represents the nematic phase, I represents the isotropic phase, pdi represents the polydispersity index and p represents the number of repeating units in the polymer. Relative molecular weights were determined by gel permeation chromatography (GPC) at 35°C using THF as solvent with polystyrene added. Variations of these examples falling within the scope of the present invention will be apparent to a person skilled in the art.

ExamplesExample 1

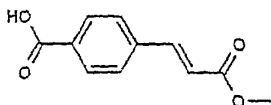
Preparation of Poly-[1-[11-[5-[4-[(*E*)-2-methoxycarbonylvinyl]benzoyloxy]-2-[6-[2-methoxy-(*E*)-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene]



A mixture of 0.70 g (0.80 mmol) (*E,E*)-5-[4-(2-methoxycarbonylvinyl)-benzoyloxy]-2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester and 1.32 mg (0.008 mmol) α,α' -azoisobutyronitrile (AIBN) in 4.0 ml dry tetrahydrofuran (THF) was degassed in a sealable tube. The tube was then sealed under argon and stirred at 60 °C for 17 h. The resulting polymer was diluted with 2.0 ml THF, precipitated into 350 ml diethyl ether and collected. The polymer was reprecipitated from THF (5.0 ml) into 400 ml methanol to yield 0.35 g (52%) Poly-[1-[11-[5-[4-[(*E*)-2-methoxycarbonylvinyl]-benzoyloxy]-2-[6-[2-methoxy-(*E*)-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-benzoyloxy]undecyloxycarbonyl]-1-methylethylene as a solid; $M_n = 7.4 \times 10^4$, $pdi = 2.12$, $T_g = 50.2$ °C, cl.p. (N/I) 68.1 °C.

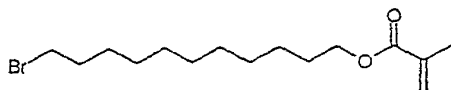
The (*E,E*)-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]-2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)-undecyl ester used as starting material was prepared in accordance with the following procedure:

Preparation of (E)-4-carboxycinnamic acid methyl ester



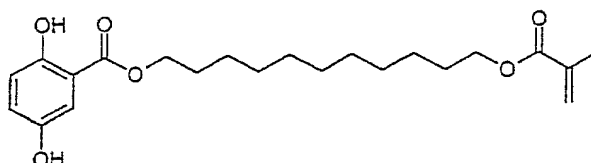
5 10.0 g (66.6 mmol) 4-carboxybenzaldehyde were dissolved in 100 ml toluene and 23.6 g (69.9 mmol) [(methoxycarbonyl)methyl]triphenylphosphorane were added. The reaction was slightly exothermic. The thick suspension was diluted with 50 ml toluene. After 18 h at room temperature the product was collected by filtration and was digested in 100 ml isopropyl alcohol at reflux temperature for 1 hour. The solid was
10 then filtered off at 0 °C, dried overnight at 45 °C under vacuum, resulting in 8.9 g (65%) (E)-4-carboxyl cinnamic acid methyl ester as white powder.

Preparation of 2-methylacrylic acid 11-bromoundecyl ester



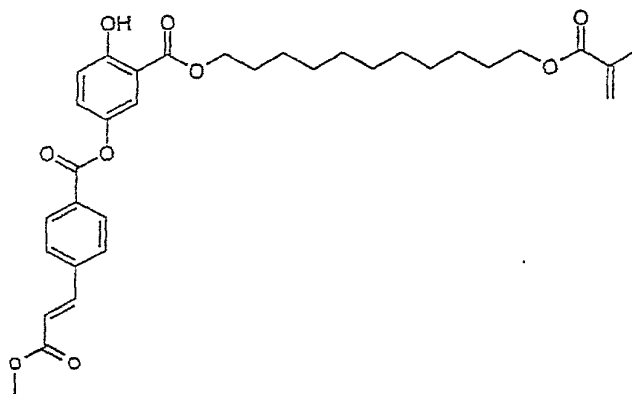
15 19.2 g (76.4 mmol) 11-bromo-1-undecanol, 7.2 g (84.1 mmol) methacrylic acid and 1.03 g (8.4 mmol) 4-dimethylaminopyridine were dissolved in 157 ml dichloromethane. The solution was subsequently cooled to 0 °C and then a solution of
20 17.4 g (84.2 mmol) dicyclohexylcarbodiimide in 80 ml dichloromethane was added dropwise at 0 °C over a period of 45 minutes. The reaction mixture was allowed to warm to room temperature, stirred for 19 hours and filtered. The filtrate was concentrated by evaporation. The residue was purified by chromatography using a silica gel column (170 g) and toluene as eluant to give 19.3 g (98.4%) 2-methylacrylic acid 11-bromoundecyl ester.
25

Preparation of 2,5-dihydroxybenzoic acid 11-(2-methylacryloyloxy)undecyl ester



- 5 8.46 g (54.9 mmol) 2,5-dihydroxybenzoic acid was suspended in 55 ml acetonitrile. 8.24 ml (54.9 mmol) 1,8-diazabicyclo[5.4.0]undec-7-ene(1,5-5) (DBU) were added dropwise over a period of 10 minutes. The reaction temperature was allowed to rise to 40 °C. The reaction mixture was cooled to room temperature and 19.3 g (60.4 mmol) 2-methacrylic acid 11-bromoundecyl ester was added and the
- 10 resulting mixture was then reflux for 17.5 hours. The reaction mixture was cooled and then extracted using diethyl ether and water; the diethyl ether was washed firstly with water, then with 1N sulfuric acid and finally with water, dried over sodium sulfate, filtered and concentrated by rotary evaporation. The residue was recrystallised twice, firstly form a mixture of ethyl acetate (17 ml) and hexane (100 ml) and secondly form
- 15 a mixture of *tert*-butyl methyl ether (50 ml) and hexane (100 ml) to give 17.1 g (79%) 2,5-dihydroxybenzoic acid 11-(2-methylacryloyloxy) ester as white crystals.

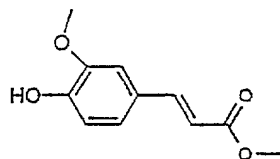
Preparation of (E)-2-Hydroxy-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoic acid 11-(2-methylacryloyloxy)undecyl ester



5

4.76 g (12.1 mmol) 2,5-dihydroxybenzoic acid 11-(2-methylacryloyloxy)ester, 2.50 g (12.1 mmol) (*E*)-4-carboxylcinnamic acid methyl ester and 0.37 g (3.0 mmol) 4-dimethylaminopyridine were dissolved in 30 ml of dichloromethane. A suspension of 2.32g (12.1 mmol) *N*-(3-dimethylaminopropyl)-*N'*-ethylcarbodiimide hydrochloride and 25 ml dichloromethane were added dropwise over a period of 1 hour. After 1.5 hour at room temperature the reaction mixture was extracted using dichloromethane and water; the dichloromethane was washed repeatedly with water, dried over sodium sulfate, filtered and concentrated by rotary evaporation. The residue was purified by chromatography using a silica gel column (240 g) using cyclohexane:ethyl acetate (9:1) as eluant to give 6.1 g (87%) (*E*)-2-hydroxy-5-[4-(2-methoxycarbonylvinyl)-benzoyloxy]benzoic acid 11-(2-methylacryloyloxy)undecyl ester as white powder, m.p. = 51 °C.

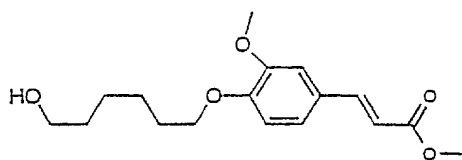
15

Preparation of (*E*)-4-Hydroxy-3-methoxycinnamic acid methyl ester

5 25 g (0.13 mol) of (*E*)-4-hydroxy-3-methoxycinnamic acid was dissolved in 180 ml of methanol, and 5 ml of concentrated sulfuric acid was added thereto. The solution was heated at reflux for 2 hours. The majority of the methanol (about 150 ml) was removed by distillation and the remaining residue was poured into 500 ml of ice-water thereby to effect precipitation of the ester, which was then purified by

10 suction filtration and washed firstly with cold water, then with a small amount of cold saturated sodium bicarbonate solution and finally with cold water and dried at 50 °C under a water-jet vacuum. The product was then purified by chromatography using silica gel (250 g) using dichloromethane:diethyl ether (19:1) as eluant to give 21.78 g of (*E*)-4-hydroxy-3-methoxycinnamic acid methyl ester in the form of a light-yellow

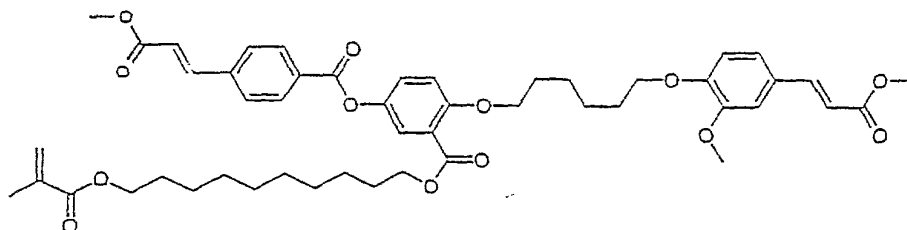
15 oil.

Preparation of (*E*)-4-(6-Hydroxyhexyloxy)-3-methoxycinnamic acid methyl ester

20 3.92 ml (25.2 mmol) of 1,6-hexanediol vinyl ether were added to a solution of 5.0 g (24.0 mmol) of (*E*)-4-hydroxy-3-methoxycinnamic acid methyl ester and 6.61 g (25.2 mmol) of triphenylphosphine in 150 ml of tetrahydrofuran. The colourless solution was subsequently cooled to 0 °C and then 11.5 ml (25.3 mmol) of a 40%

solution of azodicarboxylic acid diethyl ester in toluene were added dropwise thereto over a period of 30 minutes. The mixture was subsequently allowed to react first for 30 minutes at 0 °C and then for 22.5 hours at room temperature. 150 ml of methanol and 10 drops of concentrated sulfuric acid were then added to the reaction solution and the mixture was stirred for 1.5 hours. The reaction mixture was then extracted using ethyl acetate and water. The ethyl acetate was washed with a saturated sodium bicarbonate solution and repeatedly with saturated sodium chloride solution, dried over magnesium sulfate, filtered and concentrated by evaporation. The resulting residue was purified by column chromatography using silica gel (470 g) using toluene:ethyl acetate (1:1) as eluant. Recrystallisation from ethyl acetate:hexane (3:5) gave 6.13 g of 4-(6-hydroxyhexyloxy)-3-methoxycinnamic acid methyl ester.

Preparation of (E,E)-5-[4-(2-Methoxycarbonylvinyl)benzoyloxy]-2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)-undecyl ester

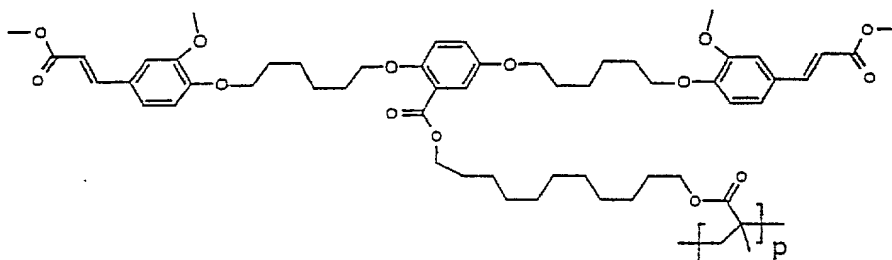


1.24 g (2.13 mmol) (E)-2-hydroxy-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoic acid 11-(2-methylacryloyloxy)undecyl ester, 0.66 g (2.13 mmol) 4-(6-hydroxyhexyloxy)-3-methoxycinnamic acid methyl ester and 0.59 g (2.24 mmol) of triphenylphosphine in 20 ml of tetrahydrofuran. The colourless solution was subsequently cooled to 0 °C and 0.98 ml (2.15 mmol) of a 40% solution of azodicarboxylic acid diethyl ester in toluene was added dropwise thereto over a period of 10 minutes. The mixture was subsequently allowed to react for 3 hours at 0 °C. The

reaction mixture was then partitioned between ethyl acetate and water; the organic phase was washed repeatedly with saturated sodium chloride solution, dried over magnesium sulfate, filtered and reduced in volume by evaporation. The resulting residue was added to a mixture of methanol and water (3:2). The resulting solid was separated from the solution by filtration and dried overnight at 45 °C under vacuum. The solid was purified by column chromatography using silica gel (150 g) and toluene:ethyl acetate (9:1) as eluant to give 1.40 g (75 %) of . (*E,E*)-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]-2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester as white powder.

Example 2

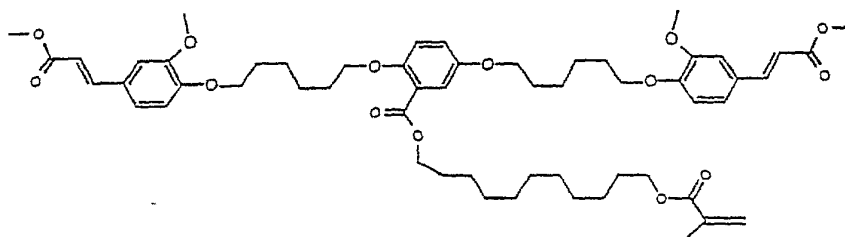
Preparation of Poly-[1-[11-[(*E,E*)-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene]



This was effected using the procedure according to Example 1 using 0.87 g (0.89 mmol) (*E,E*)-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester to give 0.25 g (30 %) poly-[1-[11-[(*E,E*)-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene] as hard solid; $M_n = 6.9 \times 10^4$, $pdi = 1.95$.

The *(E,E)*-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester used as starting material was prepared in accordance with the following procedure:

5 Preparation of *(E,E)*-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester



10 0.50 g (1.55 mmol) 2,5-dihydroxybenzoic acid 11-(2-methylacryloyloxy)ester, 0.96 g (3.10 mmol) *(E)*-4-(6-hydroxyhexyloxy)-3-methoxycinnamic acid methyl ester and 0.81 ml (3.26 mmol) of tributylphosphine were dissolved in 10 ml of tetrahydrofuran and 0.82 g (2.10 mmol) of 1,1'-(azodicarbonyl)dipiperidine were added. The mixture was allowed to react for 1 hour at room temperature. The reaction mixture was then extracted using ethyl acetate and water. The ethyl acetate was washed repeatedly with saturated sodium chloride solution, dried over magnesium sulfate, filtered and concentrated by evaporation. The resulting residue was purified by column chromatography using silica gel (150 g) using toluene:ethyl acetate (85:15) as eluant to give 0.82 g (59 %) of *(E,E)*-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester as white powder.

The following compounds were prepared in an analogous manner:

- Poly-[1-[11-[(*E,E*)-3,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)-phenoxy]oxyhexyl]benzoyloxy]undecyloxy carbonyl]-1-methylethylene]

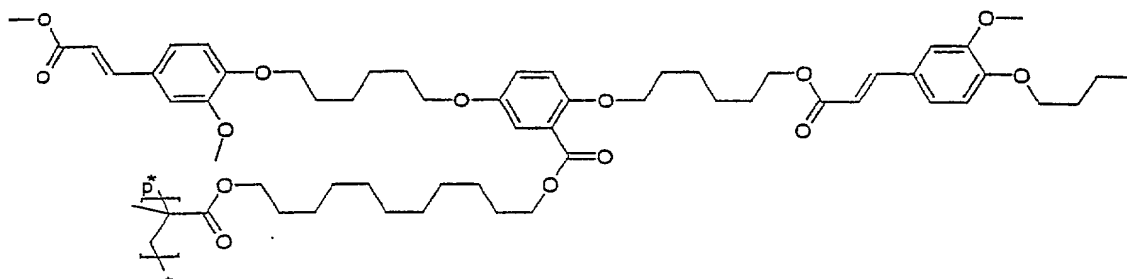
$$M_n = 1.07 \times 10^5, \text{ pdi} = 2.90$$

- Poly-[1-[11-[(*E,E*)-3,4-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)-phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene]

$$M_n = 3.59 \times 10^5, \text{ pdi} = 5.7$$

Example 3

Preparation of poly-[1-[11-[5-[6-[2-methoxy-4(*E*)-(methoxycarbonylvinyl)phenoxy]-oxyhexyl]-2-[6-[3(*E*)-(3-methoxy-4-butoxyphenyl)acryloyloxy]oxyhexyl]benzoyloxy]undecyloxy]carbonyl]-1-methylethylene]

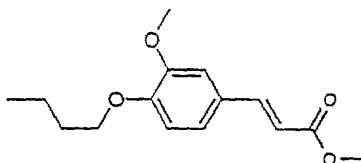


Following the procedure of Example 1 and using 0.634 g (0.624 mmol) (*E,E*)-5-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-2-[6-[3-(3-methoxy-4-butoxyphenyl)acryloyloxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)-undecyl ester and 1.0 mg (0.0062 mmol) α,α' -azoisobutyronitrile (AIBN) yielded 0.464 g (73 %) of Poly-[1-[11-[5-[6-[2-methoxy-4(*E*)-(methoxycarbonylvinyl)-phenoxy]oxyhexyl]-2-[6-[3(*E*)-(3-methoxy-4-butoxyphenyl)acryloyloxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene] as a solid; $M_n = 1.53 \times 10^5$, $pdi = 2.54$, $T_g = 17.7^\circ\text{C}$.

The (*E,E*)-5-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-2-[6-[3-(3-methoxy-4-butoxyphenyl)acryloyloxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester used as starting material was prepared in accordance with the following procedure:

5

Preparation of (*E*)-4-butoxy-3-methoxycinnamic acid methyl ester

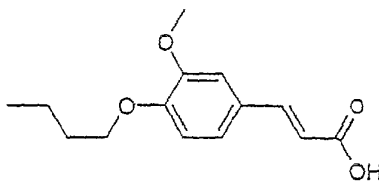


10

4.16 g (20.0 mmol) ferulic acid methyl ester was dissolved in 115 ml 2-butanone. 2.09 ml (22.0 mmol) *n*-butyl bromide and 11.06 g (80 mmol) potassium carbonate were added. The reaction suspension was then heated at reflux temperature for 20 hours. The reaction mixture was filtered. The filtrate was concentrated by evaporation. The crude product was recrystallised from 42 ml isopropyl alcohol yielded 4.85 g (92 %) (*E*)-4-butoxy-3-methoxycinnamic acid methyl ester as white crystals.

15

Preparation of (*E*)-4-butoxy-3-methoxycinnamic acid

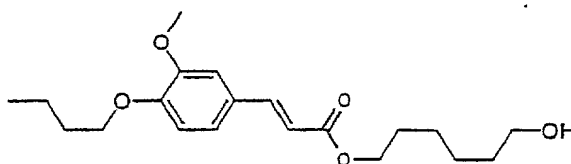


20

10 g (0.15 mol) potassium hydroxide were dissolved in a mixture of 200 ml methyl alcohol and 5 ml water. 4.85 g (18.35 mmol) (*E*)-4-butoxy-3-

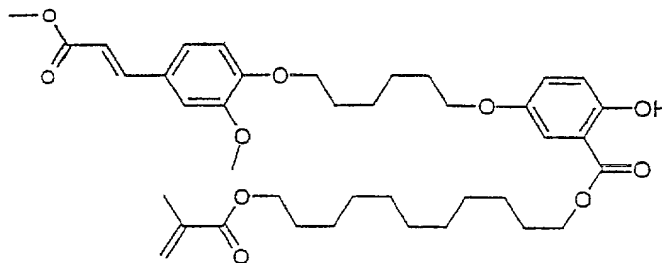
methoxycinnamic acid methyl ester was added. The reaction mixture was subsequently heated to 60°C. After 2.5 h the mixture was concentrated by evaporation. The residue was dissolved in 100 ml cold water and acidified to pH=1 with 13.5 ml hydrochloric acid 37 wt. %. The product was filtered off, washed with water and dried at 50 °C under vacuum to give 4.24 g (92 %) (*E*)-4-butoxy-3-methoxycinnamic acid as white crystals.

Preparation of (*E*)-4-butoxy-3-methoxycinnamic acid 6-hydroxyhexyl ester



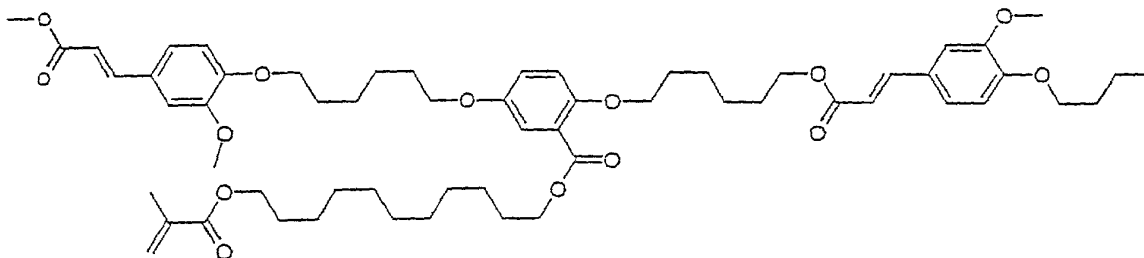
Following the procedure of Example 1 and using 1.38 g (5.50 mmol) (*E*)-4-butoxy-3-methoxycinnamic acid, 0.84 g (5.50 mmol) 1,8-diazabicyclo[5.4.0]undec-7-ene(1,5-5) (DBU) and 0.68 g (5.0 mmol) 6-chlorohexanol gave 1.39 g (79 %) (*E*)-4-butoxy-3-methoxycinnamic acid 6-hydroxyhexyl ester as colourless oil.

Preparation of (*E*)-2-Hydroxy-5-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]-oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester



1.27 g (3.24 mmol) 2,5-dihydroxybenzoic acid 11-(2-methylacryloyloxy)ester, 1.0 g (3.24 mmol) 4-(6-hydroxyhexyloxy)-3-methoxycinnamic acid methyl ester and 0.89 g (3.40 mmol) of triphenylphosphine were dissolved in 20 ml of tetrahydrofuran. The colourless solution was subsequently cooled to 0 °C and then 1.48 ml (3.40 mmol) of a 40 % solution of azodicarboxylic acid diethyl ester in toluene was added dropwise thereto over a period of 10 minutes. The mixture was subsequently allowed to react for 3 hours at 0 °C. The reaction mixture was then partitioned between ethyl acetate and water; the organic phase was washed with repeatedly with saturated sodium chloride solution, dried over magnesium sulfate, filtered and concentrated by evaporation. The residue was digest in a mixture form methanol and water 3:2. The solid was then filtered off and dried overnight at 45 °C under vacuum. Chromatography of the solid on 150 g of silica gel using toluene : ethyl acetate 1:1 yielded 1.45 g (65 %) of (*E*)-2-hydroxy-5-[6-[2-methoxy-4(methoxycarbonylvinyl)-phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy) undecyl ester as colourless oil.

Preparation of (E,E)-5-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-2-[6-[3-(3-methoxy-4-butoxyphenyl)acryloyloxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester

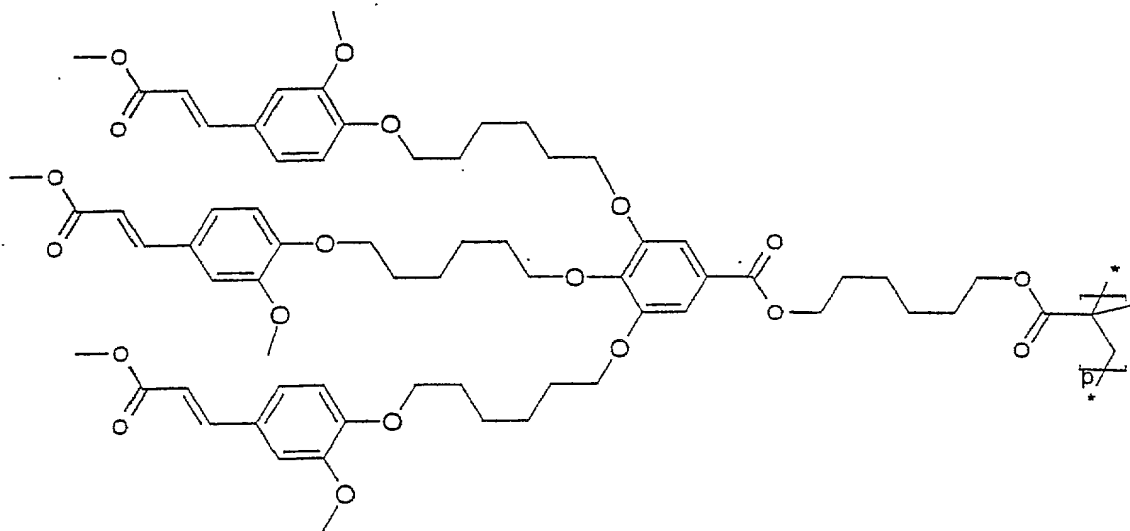


0.81 g (1.18 mmol) (*E*)-2-hydroxy-5-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy) undecyl

ester, 0.414 g (1.18 mmol) 4-(6-hydroxyhexyloxy)-3-methoxycinnamic acid methyl ester and 0.31 ml (1.24 mmol) of tributylphosphine were dissolved in 10 ml of tetrahydrofuran to which a mixture of 0.313 g (1.24 mmol) of 1,1'-(azodicarbonyl)dipiperidine and 5 ml tetrahydrofuran was subsequently added in a dropwise fashion over a period of 1 hour. The mixture was allowed to react for 18 hours at room temperature. The reaction mixture was then partitioned between ethyl acetate and water; the organic phase was washed with repeatedly with saturated sodium chloride solution, dried over magnesium sulfate, filtered and concentrated by evaporation. Chromatography of the crude product on 120 g of silica gel using toluene : ethyl acetate 4:1 and crystallisation from *tert*-butyl methyl ether yielded 0.63 g (62 %) (*E,E*)-5-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-2-[6-[3-(3-methoxy-4-butoxyphenyl)acryloyloxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester as white crystals.

15 Example 4

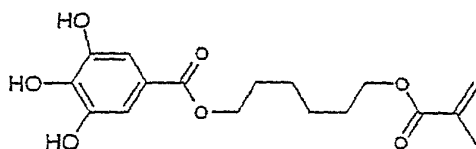
Preparation of Poly-[1-[11-[(*E,E,E*)-3,4,5-tri-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxy]carbonyl]-1-methylethylene



Preparation can be carried out analogously to Example 1 using 0.50 g (0.413 mmol) (*E,E,E*)-3,4,5-tri-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-benzoic acid 11-(2-methylacryloyloxy)undecyl ester and (0.004 mmol) α,α' -azoisobutyronitrile (AIBN) yield Poly-[1-[11-[(*E,E,E*)-3,4,5-tri-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxycarbonyl]-1-methylethylene].

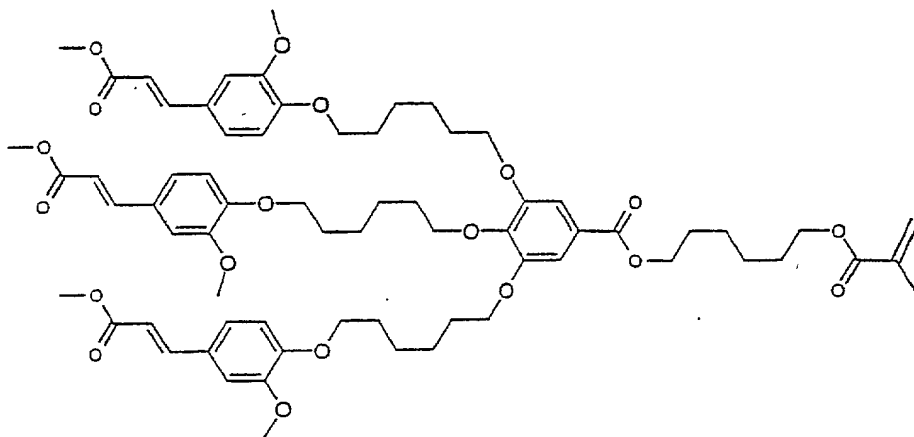
The (*E,E,E*)-3,4,5-tri-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester used as starting material was prepared in accordance with the following procedure:

Preparation of 3,4,5-dihydroxybenzoic acid 6-(2-methacryloyloxy)hexyl ester



This was effected using the procedure according to Example 1 using 6.08 g (32.3 mmol) gallic acid, 5.41 g (35.5 mmol) 1,8-diazabicyclo[5.4.0]undec-7-ene-(1,5-5) (DBU) and 6.38 g (21.5 mmol) 2-methacrylic acid 6-iodohexyl ester to give 2.43 g (33 %).

Preparation of (E,E,E)-3,4,5-tri-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester



- 5 Following the procedure of Example 1 and using 1.00 g (2.95 mmol) 3,4,5-dihydroxybenzoic acid 6-(2-methacryloyloxy)hexyl ester, 2.32 g (8.87 mmol) triphenylphosphine, 2.73 g (8.87 mmol) (E)-4-(6-hydroxyhexyloxy)3-methoxycinnamic acid methyl ester and 4.03 ml (8.87 mmol) of a 40% solution of azodicarboxylic acid diethyl ester in toluene to give 0.95 g (27 %) of (E,E,E)-
10 3,4,5-tri-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoic acid 11-(2-methylacryloyloxy)undecyl ester as colourless oil.

Example 5

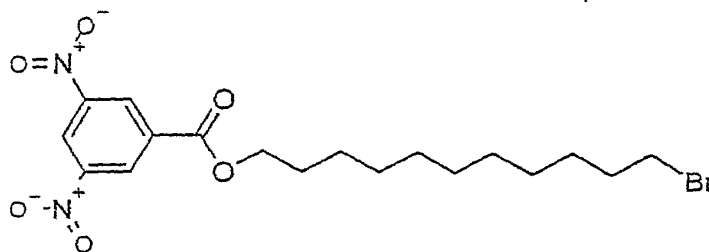
Polyimide

- 15 94.2 mg (0.4803 mmol) of 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride was added to a solution of 0.500 g (0.5336 mmol) of 3,5-diaminobenzoic acid 11-[2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]]-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoyloxy]undecyl ester in 3 ml of tetrahydrofuran. Stirring was then carried out at 0 °C for 2 hours. 10.4 mg (0.0530 mmol) of
20 1,2,3,4-cyclobutanetetracarboxylic acid dianhydride were added. The mixture was subsequently allowed to react for 69 hours at room temperature. The polymer mixture

was diluted with 3.0 ml THF, precipitated into 150 ml diethyl ether and collected. The polymer was reprecipitated from THF (10 ml) into 500 ml water to yield, after drying at room temperature under vacuum, polyamic acid.

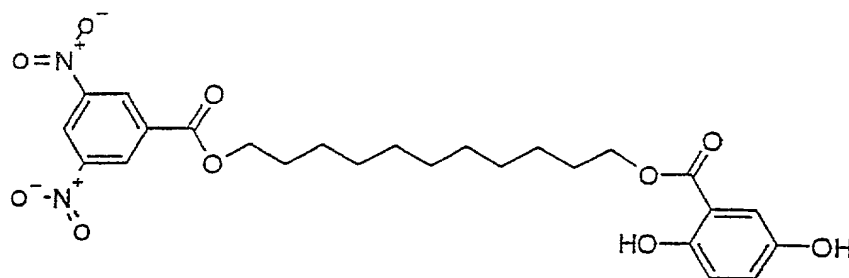
The 3,5-diaminobenzoic acid 11-[2-[6-[2-methoxy-4-(methoxycarbonylvinyl)-phenoxy]oxyhexyl]]-5-[4-(2-methoxycarbonyl-vinyl)benzoyloxy]benzoyloxy]undecyl ester used as starting material was prepared in accordance with the following procedure:

3,5-Dinitrobenzoic acid 11-bromoundecyl ester



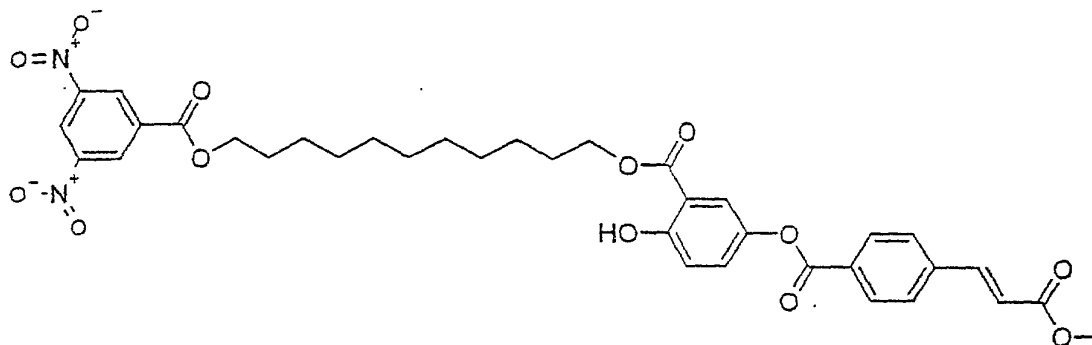
11.4 g (45.4 mmol) 11-bromo-1-undecanol, 11.0 g (47.7 mmol) 3,5-dinitrobenzoyl chloride and 54 mg 4-dimethylaminopyridine were dissolved in 94 ml dichloromethane. The solution was subsequently cooled to 0 °C and then 18.3 ml (227 mmol) pyridine was added dropwise, in the course of 25 minutes. After 4.5 hours at 0 °C the reaction mixture was partitioned between dichloromethane and water. The organic phase was washed repeatedly with water, dried over sodium sulfate, filtered and concentrated by rotary evaporation. Chromatography of the residue on 200 g silica gel using toluene yielded 18.1 g (90 %) 3,5-dinitrobenzoic acid 11-bromoundecyl ester as yellow powder.

3,5-Dinitrobenzoic acid 11-[2,5-dihydroxybenzoyloxy]undecyl ester



2.78 g (18.0 mmol) 2,5-dihydroxybenzoic acid was dissolved in 36 ml
 5 dimethylformamide. 2.96 ml (19.8 mmol) 1,8-diazabicyclo[5.4.0]undec-7-ene(1,5-5)
 (DBU) was added dropwise in the course of 15 minutes. The reaction temperature rose
 to 30 °C and 8.83 g (19.8 mmol) 3,5-dinitrobenzoic acid 11-bromoundecyl ester was
 subsequently added in one portion. The mixture was then heated at 80 °C for 2 hours.
 The reaction mixture was cooled and then partitioned between dichloromethane and
 10 1N hydrochloric acid; the organic phase was washed twice with water, dried over
 sodium sulfate, filtered and concentrated by rotary evaporation. Chromatography of
 the residue on 200 g silica gel using Toluene yielded 5.28 g (57%) 3,5-dinitrobenzoic
 acid 11-[2,5-dihydroxybenzoyloxy]undecyl ester as yellow powder.

3,5-Dinitrobenzoic acid 11-[2-hydroxy-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]-benzoyloxy]undecyl ester



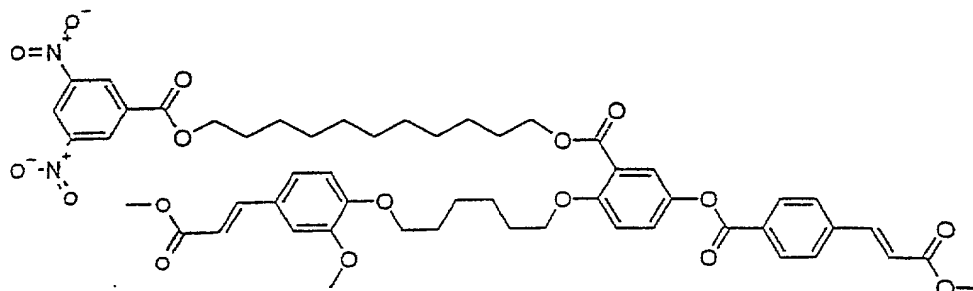
5

2.50 g (4.82 mmol) 3,5-Dinitrobenzoic acid 11-[2,5-dihydroxybenzoyloxy]undecyl ester, 0.99 g (4.82 mmol) (*E*)-4-carboxyl cinnamic acid methyl ester and 0.15 g (1.20 mmol) 4-dimethylaminopyridine were dissolved in 12 ml of dichloromethane.

10 A suspension of 0.92 g (4.82 mmol) *N*-(3-dimethylaminopropyl)-*N'*-ethylcarbodiimide hydrochloride and 10 ml dichloromethane were added dropwise in the course of 45 minutes. After 3 hours at room temperature the reaction mixture was partitioned between dichloromethane and water; the organic phase was washed repeatedly with water, dried over sodium sulfate, filtered and concentrated by rotary evaporation.

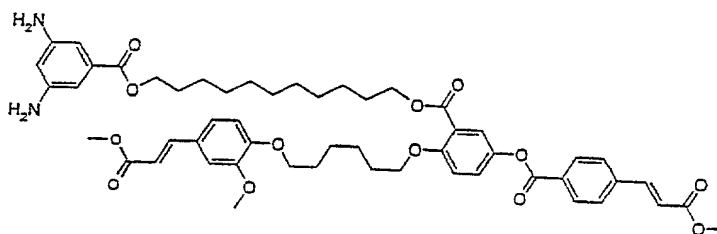
15 Chromatography of the residue on 100 g silica gel using toluene yielded 2.63 g (77%) 3,5-dinitrobenzoic acid 11-[2-hydroxy 5-[4-(2-methoxycarbonyl-vinyl)benzoyloxy]benzoyloxy]undecyl ester as yellow powder.

3,5-Dinitrobenzoic acid 11-[2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]-oxyhexyl]-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoyloxy]undecyl ester



1.00 g (1.41 mmol) 3,5-dinitrobenzoic acid 11-[2-hydroxy-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoyloxy]undecyl ester, 0.436 g (1.41 mmol) (*E*)-4-(6-hydroxyhexyloxy)-3-methoxycinnamic acid methyl ester and 0.37 ml (1.48 mmol) of tributylphosphine were dissolved in 10 ml of tetrahydrofuran and 0.37 g (1.48 mmol) of 1,1'-(azodicarbonyl)dipiperidine is added. The mixture was allowed to react for 1 hour at room temperature. The reaction mixture was then partitioned between ethyl acetate and water; the organic phase was washed with repeatedly with saturated sodium chloride solution, dried over magnesium sulfate, filtered and concentrated by evaporation. Chromatography of the solid on of silica gel yield 3,5-Dinitrobenzoic acid 11-[2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoyloxy]undecyl ester.

3,5-Diaminobenzoic acid 11-[2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]-oxyhexyl]-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoyloxy]undecyl ester

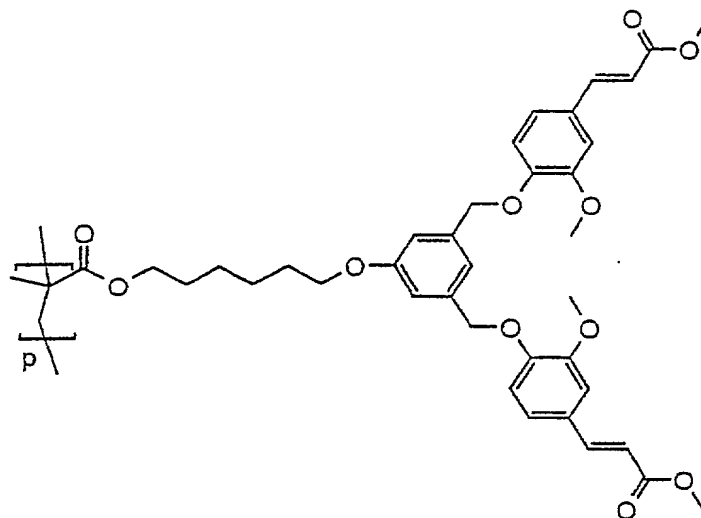


5

0.997 g (1.00 mmol) 3,5-Dinitrobenzoic acid 11-[2-[6-[2-methoxy-4-(methoxycarbonylvinyl) phenoxy]oxyhexyl]-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoyloxy]undecyl ester and 0g (4.00 mmol) of ammonium chloride were suspended in 15 ml of a mixture consisting of methanol : water 9:1. 1.31 g (20.0 mmol) of zinc was then added in one portions. After 0.5 hour at room temperature 20 ml of a mixture consisting of methanol : water 9:1 is added to the thick suspension. After a further 21 hours the reaction suspension was partitioned between dichloromethane and water. The resulting suspension was filtered, the organic phase was washed with a saturated sodium bicarbonate solution and repeatedly with water. The organic phase was then dried over sodium sulfate, filtered and concentrated by evaporation. Chromatography of the residue on silica yield 3,5-Diaminobenzoic acid 11-[2-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]-5-[4-(2-methoxycarbonylvinyl)benzoyloxy]benzoyloxy]undecyl ester.

Example 6

Preparation of Poly-[1-[6-[3,5-bis[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyloxycarbonyl]-1-methyl-ethylene]



5

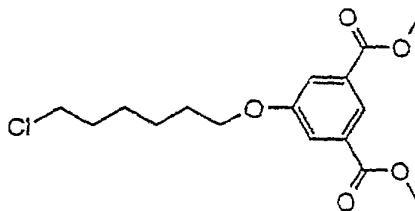
A solution of 0.46 g (0.63 mmol) 2-methylacrylic acid 6-[3,5-bis[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyl ester and 1.0 mg (0.0063 mmol) α,α' -azoisobutyronitrile (AIBN) in 1.6 ml dry tetrahydrofuran (THF) was degassed in a Schlenk tube and sealed under argon. The mixture was stirred at 55 °C for 15 h. The resulting polymer was precipitated into 500 ml methanol and collected. The polymer was reprecipitated from 3.0 ml THF into 500 ml diethylether at 0 °C to yield 0.2 g (46%) Poly-[1-[6-[3,5-bis[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyloxycarbonyl]-1-methyl-ethylene] as a white solid.

15

The 2-methylacrylic acid 6-[3,5-bis[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyl ester used as starting material was prepared in accordance with the following procedure:

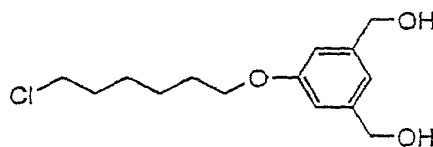
20

Preparation of 5-[(6-Chlorohexyl)oxy]isophthalic acid dimethyl ester



13.02 g (95.3 mmol) of 6-chlorohexanol were added to a solution of 20.04 g (95.3 mmol) of 5-hydroxyisophthalic acid dimethyl ester and 47.4 g (180.5 mmol) of triphenylphosphine in 300 ml of tetrahydrofuran. The colourless solution was cooled to 0 °C and then 79 ml (180.5 mmol) of a 40% solution of azodicarboxylic acid diethyl ester in toluene were added dropwise over a period of 2 hours and 45 minutes. The mixture was allowed to react for another 30 minutes at 0 °C and then for 22 hours at room temperature. The reaction mixture was concentrated by evaporation under vacuum. The residue was filtered through silica gel using dichloromethane:diethyl-ether (85:15) as eluent. The filtrate was concentrated in vacuum and further purified by column chromatography on silica gel with hexane:ethyl acetate (9:1) to give 17.12 g (55%) of 5-[(6-chlorohexyl)oxy]isophthalic acid dimethyl ester as a white solid.

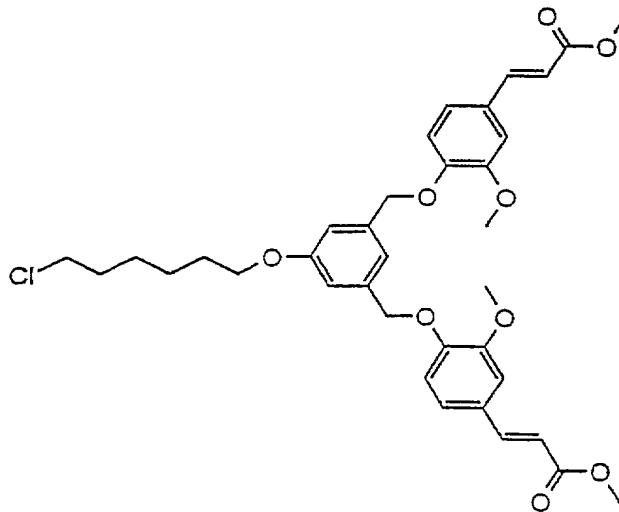
Preparation of [3-[(6-Chlorohexyl)oxy]-5-[hydroxymethyl]phenyl]methanol



6.5 g (19.8 mmol) 5-[(6-chlorohexyl)oxy]isophthalic acid dimethyl ester were dissolved in 65 ml tetrahydrofuran in an atmosphere of argon. The colourless solution was cooled to -25 °C and then 42 ml (41.6 mmol) of a 1 molar solution of lithiumaluminiumhydride in tetrahydrofuran were added dropwise over a period of 30

minutes. The reaction mixture was carefully quenched in sequence first with 5 ml of methanol and then with 50 ml of a 1 molar solution of HCl in water. The suspension was stirred for 1 hour at room temperature and subsequently filtered through speedex. The solid residue was washed carefully with 100 ml of tert.-butyl-methylether. The organic phase of the filtrate was separated, washed with 100 ml of a saturated sodium bicarbonate solution and 100 ml of water, dried over magnesium sulfate, filtered and concentrated in vacuum. Column chromatography of the crude product on 250 g of silica gel using dichloromethane:methanol (9:1) as eluent gave 5 g (93%) of [3-[(6-chlorohexyl)oxy]-5-[hydroxymethyl]phenyl]methanol as a white solid.

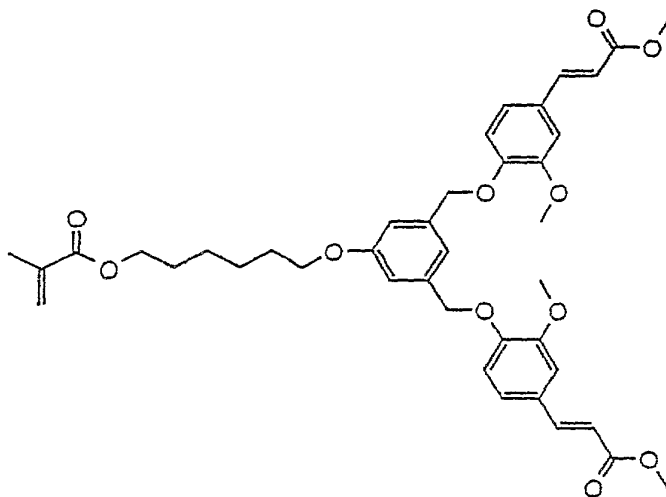
Preparation of (E)-3-[4-[[3-[(6-Chlorohexyl)oxy]-5-[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]benzyl]oxy]-3-methoxyphenyl]acrylic acid methyl ester



1.5 g (5.5 mmol) [3-[(6-chlorohexyl)oxy]-5-[hydroxymethyl]phenyl] methanol were added to a solution of 2.86 g (13.7 mmol) (E)-4-hydroxy-3-methoxycinnamic acid methyl ester (see Example 1) and 3.61 g (13.7 mmol) triphenylphosphine in 20 ml tetrahydrofuran. The colourless solution was cooled to 0 °C and then 6 g (13.7 mmol) of a 40% solution of azodicarboxylic acid diethyl ester in toluene were added

dropwise over a period of 2 hours. The mixture was allowed to react for 2 hours at 0 °C and then for 22 hours at room temperature. The reaction mixture was concentrated in vacuum. The solid residue was dissolved in 50 ml of dichloromethane, washed with 30 ml of an 1 M HCl solution, 50 ml of a saturated sodium bicarbonate solution and repeatedly with water, dried over magnesium sulfate, filtered and concentrated by vacuum evaporation. The semicrystalline residue was dissolved in 15 ml methanol at 60 °C. The final product crystallized on standing at -20 °C overnight. Recrystallization from 38 ml of methanol gave 3.5 g (96%) of (*E*)-3-[4-[[3-[(6-chlorohexyl)oxy]-5-[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]-benzyl]oxy]-3-methoxyphenyl]acrylic acid methyl ester as white crystals.

Preparation of 2-Methylacrylic acid 6-[3,5-bis[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyl ester

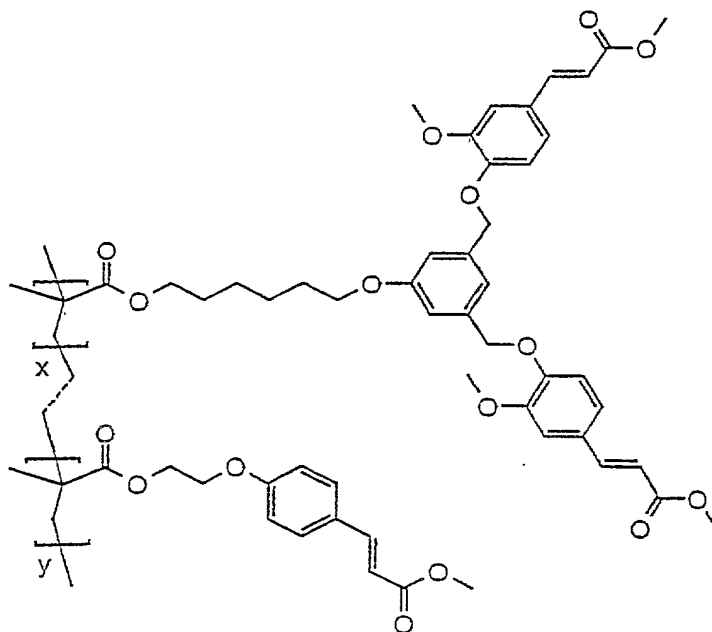


382 mg (2.5 mmol) 1,8-diazabicyclo[5.4.0]undec-7-ene(1,5-5) (DBU) in 4 ml N,N-dimethylformamide were added dropwise to a solution of 200 mg (2.3 mmol) 2-methylacrylic acid in 8 ml N,N-dimethylformamide over a period of 35 minutes. After addition of 4 mg phenothiazine, 1.5 g (2.3 mmol) (*E*)-3-[4-[[3-[(6-chlorohexyl)oxy]-5-

[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]benzyl]oxy]-3-methoxyphenyl]acrylic acid methyl ester and 84 mg tetrabutylammonium iodide the resulting mixture was stirred at 80 °C for 25 hours. The reaction mixture was poured into 50 ml of an ice cold 1 M HCl solution. The aqueous phase was extracted three times with 50 ml ethyl acetate. The organic phase was washed with 50 ml of a saturated sodium bicarbonate solution and repeatedly with water, dried over magnesium sulfate, filtered and concentrated by vacuum evaporation. Purification of the residue by column chromatography on 275 g silica gel using hexane:ethyl acetate (1:1) as eluent yielded 1.46 g (90%) of 2-methylacrylic acid 6-[3,5-bis[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyl ester as yellowish oil.

Example 7

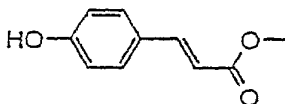
Preparation of Poly-[1-[6-[3,5-bis[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyloxycarbonyl]-1-methyl-ethylene-co-1-[2-[4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]ethoxycarbonyl]-1-methyl-ethylene] (1:1)



A solution of 0.50 g (0.71 mmol) 2-methylacrylic acid 6-[3,5-bis[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyl ester, 0.21 g (0.71 mmol) 2-methylacrylic acid 2-[4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]ethyl ester and 2.3 mg (0.014 mmol) α,α' -azoisobutyronitrile (AIBN) in 2.8 ml dry tetrahydrofuran (THF) was degassed in a Schlenk tube and sealed under argon. The mixture was stirred at 55 °C for 21 h. The resulting polymer was precipitated into 700 ml diethylether and collected. The polymer was reprecipitated from 5 ml THF into 500 ml diethylether to yield 0.58 g (82%) Poly-[1-[6-[3,5-bis[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]phenoxy]hexyloxycarbonyl]-1-methyl-ethylene-co-1-[2-[4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]ethoxycarbonyl]-1-methyl-ethylene] (1:1) as a white solid.

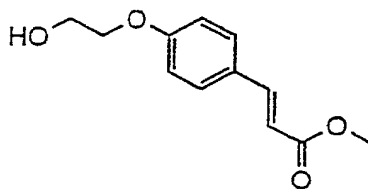
The 2-methylacrylic acid 2-[4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]ethyl ester used as comonomer was prepared according to the following procedure:

Preparation of 3-(4-Hydroxyphenyl)acrylic acid methyl ester



51.2 g (312 mmol) of p-coumaric acid were dissolved in 330 ml of methanol and treated with 10 ml of concentrated sulphuric acid. The solution was heated under reflux for 2 hours. Subsequently the majority of the methanol (about 200 ml) was distilled off and the residue remaining behind was poured into 1.3 l of ice-water. The separated ester was filtered off under suction and washed in succession with cold water, with a small amount of cold NaHCO_3 solution and again with cold water. Drying at 50 °C in a water-jet vacuum gave 51.1 g of 3-(4-hydroxyphenyl)acrylic acid methyl ester in the form of a light brownish coloured powder.

Preparation of (*E*)-3-[4-[2-Hydroxyethoxy]phenyl]acrylic acid methyl ester

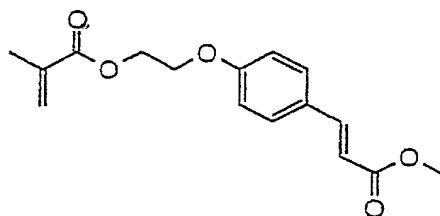


5

30 g (168 mmol) of 3-(4-hydroxyphenyl)acrylic acid methyl ester, 29 g (210 mmol) of anhydrous K_2CO_3 and a spatula tip of KI were placed in 200 ml of dimethylformamide. 14.91 g (185 mmol) of 2-chloroethanol were added dropwise at 85 °C within 5 minutes while stirring. The batch was stirred at 85 °C for a further 3 days. Subsequently, the salts were filtered off and the filtrate was concentrated to dryness in a water-jet vacuum. 16.1 g of (*E*)-3-[4-[2-hydroxyethoxy]phenyl]acrylic acid methyl ester were obtained in the form of white crystals after recrystallization from i-propanol.

10

15 Preparation of 2-Methylacrylic acid 2-[4-[*E*)-2-(methoxycarbonyl)vinyl]phenoxy]-ethyl ester



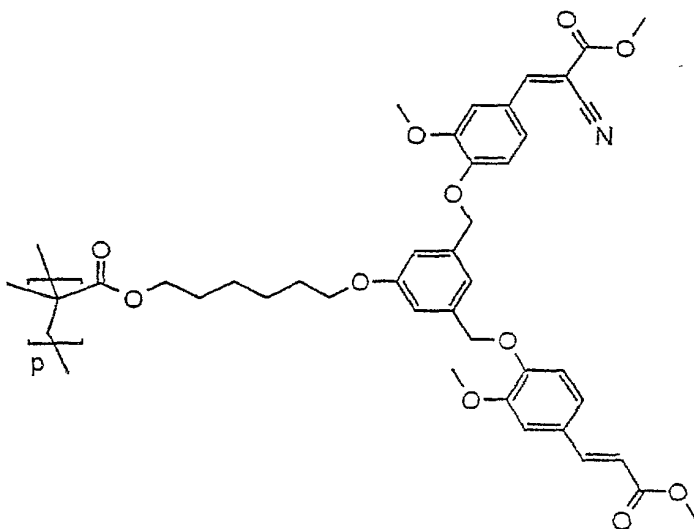
20

2.56 g (30 mmol) of methacrylic acid in 10 ml of THF were slowly added dropwise to a solution of 6 g (27 mmol) of (*E*)-3-[4-[2-hydroxyethoxy]phenyl]acrylic acid methyl ester, 5.85 g (28.3 mmol) of *N,N'*-dicyclohexylcarbodiimide (DCC) and 0.37 g (3 mmol) of 4-dimethylamino-pyridine in 80 ml of tetrahydrofuran (THF). The batch was stirred at room temperature overnight. In order to complete the reaction

there were added firstly a further 1.46 g (7.1 mmol) of DCC and, after stirring for one hour, a further 0.5 g (5.9 mmol) of methacrylic acid. The batch was stirred for a further 24 hours, filtered and the filtrate was extracted 3 times each time with 200 ml of 5% acetic acid and 200 ml of water. The ether phase was dried over Na₂SO₄,
 5 evaporated and the residue was recrystallized from cyclohexane. Subsequently, the still slightly impure product was filtered over a thin silica gel layer (eluent: diethyl ether/hexane = 1:1). This gave 8.3 g of 2-methylacrylic acid 2-[4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]ethyl ester as a white powder with a melting point of 81-82 °C and an absorption maximum of λ_{max} (in CH₂Cl₂) = 306.5 nm (ϵ = 23675 l/mol
 10 cm).

Example 8

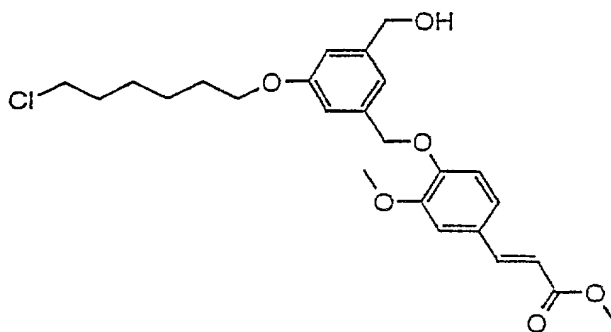
Preparation of Poly-[1-[6-[3-[2-methoxy-4-[2-cyano-(*E*)-2-(methoxycarbonyl)vinyl]-phenoxy]methyl]-5-[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]-15 phenoxy]hexyloxy]carbonyl]-1-methyl-ethylene]



A solution of 0.35 g (0.48 mmol) 2-methylacrylic acid 6-[3-[2-methoxy-4-[2-cyano-(*E*)-2-(methoxycarbonyl)vinyl]phenoxy-methyl]-5-[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy-methyl]phenoxy]hexyl ester and 0.78 mg (0.0048 mmol) α,α' -azoisobutyronitrile (AIBN) in 1.2 ml dry tetrahydrofuran (THF) was degassed in a Schlenk tube and sealed under argon. The mixture was stirred at 55 °C for 16.5 h. The resulting polymer was precipitated into 350 ml methanol and collected. The polymer was reprecipitated from 3 ml THF into 350 ml methanol to yield 0.11 g (31%) Poly-[1-[6-[3-[2-methoxy-4-[2-cyano-(*E*)-2-(methoxycarbonyl)vinyl]phenoxy-methyl]-5-[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy-methyl]phenoxy]-hexyloxycarbonyl]-1-methyl-ethylene] as a yellow solid.

The 2-methylacrylic acid 6-[3-[2-methoxy-4-[2-cyano-(*E*)-2-(methoxycarbonyl)vinyl]phenoxy-methyl]-5-[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy-methyl]phenoxy]hexyl ester used as starting material was prepared in accordance with the following procedure:

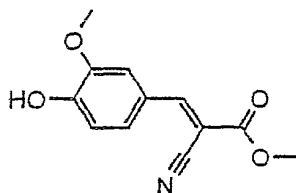
Preparation of (*E*)-3-[4-[3-[(6-Chlorohexyl)oxy]-5-[hydroxymethyl]benzyloxy]-3-methoxyphenyl]acrylic acid methyl ester



2.7 g (10 mmol) [3-[(6-chlorohexyl)oxy]-5-[hydroxymethyl]phenyl] methanol (Example 6) were added to a solution of 2.6 g (12.5 mmol) (*E*)-4-hydroxy-3-methoxy-

cinnamic acid methyl ester (Example 1) and 3.38 g (12.5 mmol) triphenylphosphine in 40 ml tetrahydrofuran. The colourless solution was cooled to 0 °C and then 5.2 ml (12.5 mmol) of a 40% solution of azodicarboxylic acid diethyl ester in toluene were added dropwise over a period of 2 hours. The mixture was allowed to react for 2 hours
5 at 0 °C and then for 15 hours at room temperature. The reaction mixture was poured into an ice cold 1 M solution of HCl. The aqueous phase was extracted three times with a mixture of tert.-butyl-methylether/ethyl acetate. The organic phase was washed with a saturated sodium bicarbonate solution and repeatedly with water, dried over magnesium sulfate, filtered and concentrated in vacuum. Purification of the residue by
10 column chromatography on silica gel using hexane:ethyl acetate (1:1) as eluent yielded 2.2 g (48%) of (*E*)-3-[4-[3-[(6-chlorohexyl)oxy]-5-[hydroxymethyl]benzyl-oxy]-3-methoxyphenyl]acrylic acid methyl ester as an oily product that solidified on standing.

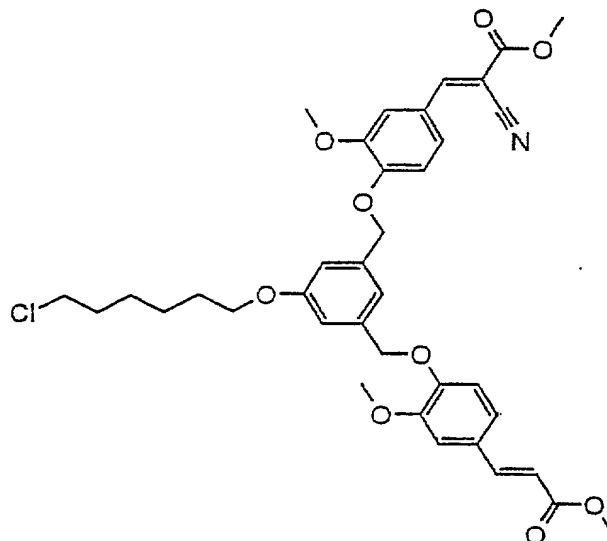
15 Preparation of 2-Cyano-(*E*)-3-(4-hydroxy-3-methoxyphenyl)acrylic acid methyl ester



0.2 ml of piperidine were added carefully to a solution of 4.6 g (30 mmol) of
20 vanillin, 3.4 ml (30.3 mmol) of cyanoacetic acid methyl ester and 0.5 ml of ethanol held at 45 °C. The product started to precipitate on cooling to room temperature. The cold reaction mixture was diluted with 8 ml of ethanol, filtered and washed with cold ethanol to give 6.5 g (93%) of 2-cyano-(*E*)-3-(4-hydroxy-3-methoxyphenyl)acrylic acid methyl ester in form of yellow crystals.

25

Preparation of (E)-3-[4-[[3-[(6-Chlorohexyl)oxy]-5-[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]benzyl]oxy]-3-methoxyphenyl]-2-cyanoacrylic acid methyl ester



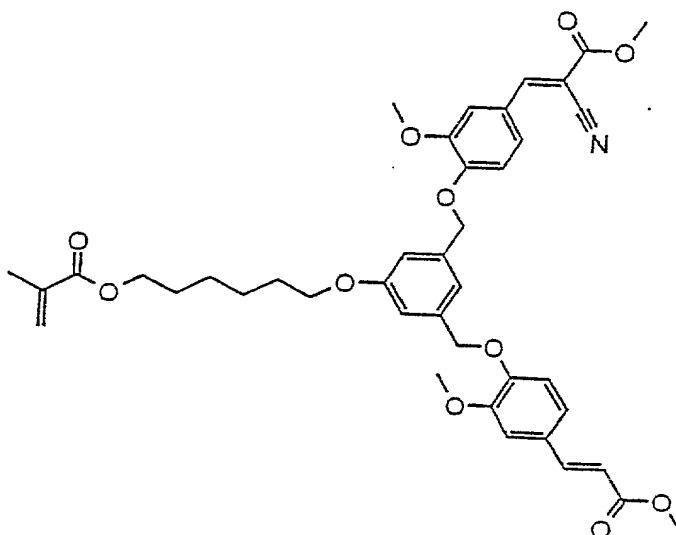
5

1 g (2.2 mmol) (E)-3-[4-[3-[(6-chlorohexyl)oxy]-5-[hydroxymethyl]benzyl-oxy]-3-methoxyphenyl]acrylic acid methyl ester was added to a solution of 0.62 g (2.7 mmol) 2-cyano-(E)-3-(4-hydroxy-3-methoxyphenyl)acrylic acid methyl ester and 0.73 g (2.7 mmol) triphenylphosphine in 10 ml tetrahydrofuran. The solution was cooled to 0 °C and then 1.1 ml (2.7 mmol) of a 40% solution of azodicarboxylic acid diethyl ester in toluene were added dropwise. The mixture was allowed to react for 1 hour at 0 °C and then it was poured into an ice cold 1 M solution of HCl. The mixture was extracted three times with tert.-butyl-methylether/ethyl acetate. The organic phase was washed with a saturated sodium bicarbonate solution and repeatedly with water, dried over magnesium sulfate, filtered and concentrated in vacuum. The solid residue was first treated with diethylether, filtered and then it was recrystallized twice from tert.-butyl-methylether to give 1.1 g (79%) of (E)-3-[4-[[3-[(6-Chlorohexyl)oxy]-5-[[2-

15

methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]benzyl]oxy]-3-methoxyphenyl]-2-cyanoacrylic acid methyl ester in form of slightly yellow crystals.

Preparation of 2-Methylacrylic acid 6-[3-[2-methoxy-4-[2-cyano-(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]-5-[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]-phenoxy]methyl]phenyl]hexyl ester.

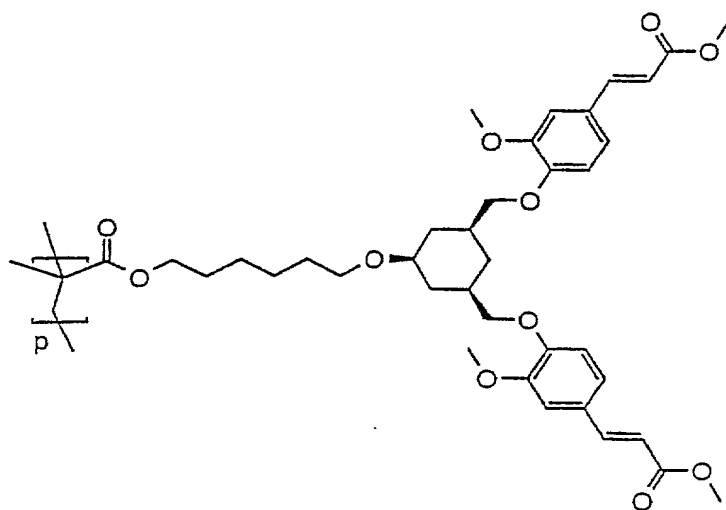


0.32 ml (2 mmol) 1,8-diazabicyclo[5.4.0]undec-7-ene(1,5-5) (DBU) in 2 ml N,N-dimethylformamide (DMF) were added dropwise to a solution of 0.2 ml (1.9 mmol) 2-methylacrylic acid in 2 ml DMF. After addition of 2 mg phenothiazine, 60 mg tetrabutylammonium iodide and 1.1 g (1.6 mmol) (*E*)-3-[4-[[3-[(6-chlorohexyl)-oxy]-5-[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]benzyl]oxy]-3-methoxyphenyl]-2-cyanoacrylic acid methyl ester in 7 ml DMF the resulting mixture was stirred at 80 °C for 20 hours. The reaction mixture was poured into an ice cold 1 M HCl solution. The aqueous phase was extracted three times with ethyl acetate. The organic phase was washed with a saturated sodium bicarbonate solution and repeatedly with water, dried over magnesium sulfate, filtered and concentrated by

vacuum evaporation. The crude product was filtered through a thin silica gel layer (eluent: ethyl acetate). The filtrate was evaporated to dryness and recrystallized from ethanol to yield 0.35 g (30%) of 2-methylacrylic acid 6-[3-[2-methoxy-4-[2-cyano-
 5 carbonyl]vinyl]phenoxy]methyl]phenoxy]hexyl ester as slightly yellow crystals.

Example 9

Preparation of Poly-[1-[6-[3,5-bis[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]-
 phenoxy]methyl]cyclohexyloxy]hexyloxycarbonyl]-1-methyl-ethylene]

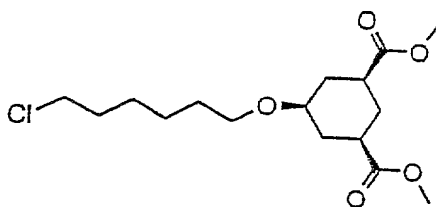


A solution of 0.5 g (0.7 mmol) 2-methylacrylic acid 6-[3,5-bis[[2-methoxy-4-
 15 [(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]cyclohexyloxy]hexyl ester and 1.1
 mg (0.007 mmol) α,α' -azoisobutyronitrile (AIBN) in 1.7 ml dry tetrahydrofuran
 (THF) was degassed in a Schlenk tube and sealed under argon. The mixture was
 stirred at 55 °C for 38 h. The resulting polymer was precipitated into 500 ml
 diethylether and collected. The polymer was reprecipitated from 3.5 ml THF into 500
 ml diethylether to yield 0.29 g (58%) Poly-[1-[6-[3,5-bis[[2-methoxy-4-[(E)-2-

(methoxycarbonyl)vinyl]phenoxy)methyl]cyclohexyloxy]hexyloxycarbonyl]-
1-methyl-ethylene] as a white solid.

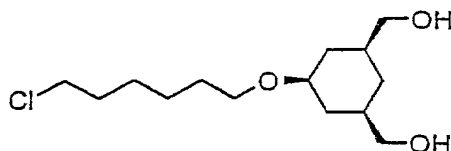
The 2-methylacrylic acid 6-[3,5-bis[[2-methoxy-4-[(*E*)-2-(methoxycarbonyl)-
5 vinyl]phenoxy)methyl]cyclohexyloxy]hexyl ester used as starting material was
prepared in accordance with the following procedure:

Preparation of 5-[(6-Chlorohexyl)oxy]-1,3-cyclohexanedicarboxylic acid dimethyl
ester



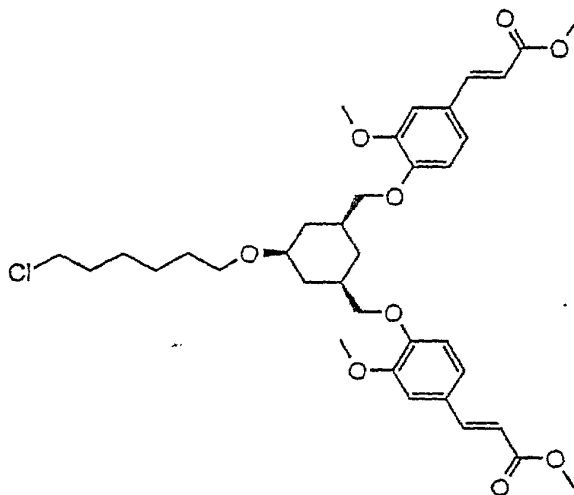
10 7.5 g (22.8 mmol) 5-[(6-chlorohexyl)oxy]isophthalic acid dimethyl ester
(example 6) were dissolved in 75 ml ethyl acetate and charged into a steel autoclave.
After addition of 3 g of a hydrogenation catalyst (5% Rh/Alox, Johnson Matthey
Company) the vessel was closed and the compound was hydrogenated at a reaction
15 temperature of 70 °C using a constant hydrogen pressure of 18 bar until no further
hydrogen uptake was observed (approx. 4 hours). The reaction suspension was cooled
to room temperature, filtered and concentrated under vacuum to dryness. The crude
product was essentially a cis/trans mixture of 5-[(6-chlorohexyl)oxy]-1,3-cyclo-
hexanedicarboxylic acid dimethyl ester. The pure cis-compound was isolated as a
20 colourless oil by chromatography on silica gel using hexane/ethyl acetate (4:1) as
eluent. (yield: 6.1 g, 80%).

Preparation of [3-[(6-Chlorohexyl)oxy]-5-[hydroxymethyl]cyclohexyl]methanol



5 6.05 g (17.5 mmol) cis-5-[(6-chlorohexyl)oxy]-1,3-cyclohexanedicarboxylic acid dimethyl ester were dissolved in 60 ml tetrahydrofuran in an atmosphere of argon. The colourless solution was cooled to -25 °C and then 37 ml (36.8 mmol) of a 1 molar solution of lithiumaluminiumhydride in tetrahydrofuran were added dropwise over a period of 45 minutes. The reaction mixture was carefully quenched in sequence
10 first with 5 ml of methanol and then with 50 ml of a 1 molar solution of HCl in water. The suspension was stirred for 1 hour at room temperature and subsequently filtered through speedex. The solid residue was washed carefully with 100 ml of tert.-butylmethylether. The organic phase of the filtrate was separated, washed with 100 ml of a saturated sodium bicarbonate solution and 100 ml of water, dried over magnesium
15 sulfate, filtered and concentrated in vacuum. Column chromatography of the crude product on 280 g of silica gel using dichloromethane:methanol (19:1) as eluent gave 4.6 g (94%) of [3-[(6-chlorohexyl)oxy]-5-[hydroxymethyl]cyclohexyl]methanol as a colourless oil.

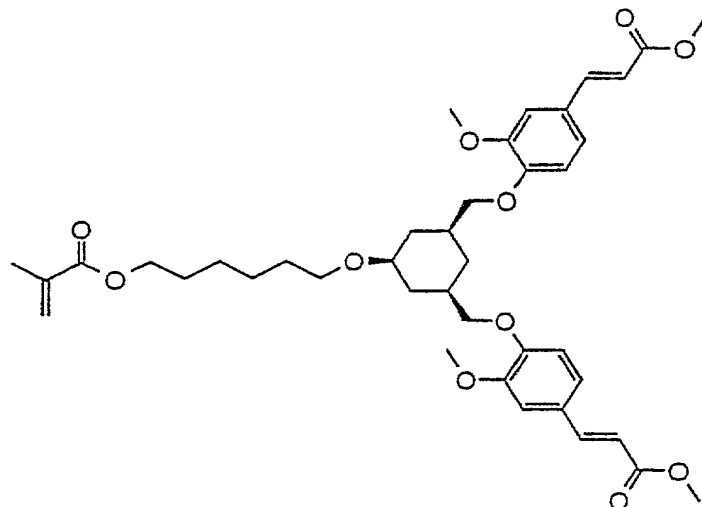
Preparation of (E)-3-[4-[[3-[(6-Chlorohexyl)oxy]-5-[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]cyclohexyl]methoxy]-3-methoxyphenyl]acrylic acid methyl ester



2.25 g (7.9 mmol) [3-[(6-chlorohexyl)oxy]-5-[hydroxymethyl]cyclohexyl]-methanol, 4.12 g (19.8 mmol) (E)-4-hydroxy-3-methoxycinnamic acid methyl ester (Example 1) and 5.24 g (19.8 mmol) triphenylphosphine were solved in 25 ml tetrahydrofuran. The solution was cooled to 0 °C and then 8.64 g (19.8 mmol) of a 40% solution of azodicarboxylic acid diethyl ester in toluene were added dropwise over a period of 2 hours. The mixture was allowed to react for 2 hours at 0 °C and then for 24 hours at room temperature. The reaction mixture was concentrated in vacuum. The solid residue was dissolved in 50 ml of dichloromethane, washed with 50 ml of an 1 M HCl solution, 50 ml of a saturated sodium bicarbonate solution and repeatedly with water, dried over magnesium sulfate, filtered and concentrated by vacuum evaporation. The crude product was crystallized from methanol at -20 °C. Further purification of the solid residue by chromatography on silica gel using dichloromethane/ethyl acetate (19:1) as eluent gave 2.6 g (50%) of (E)-3-[4-[[3-[(6-chlorohexyl)oxy]-5-[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]-

cyclohexyl]methoxy]-3-methoxyphenyl]acrylic acid methyl ester as a colourless highly viscous oil that solidified on standing.

Preparation of 2-Methylacrylic acid 6-[3.5-bis[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]cyclohexyloxy]hexyl ester



786 mg (5.16 mmol) 1,8-diazabicyclo[5.4.0]undec-7-ene(1,5-5) (DBU) in 5 ml N,N-dimethylformamide were added dropwise to a solution of 413 mg (4.8 mmol) 2-methylacrylic acid in 5 ml N,N-dimethylformamide over a period of 30 minutes. After addition of 4 mg phenothiazine, 146 mg tetrabutylammonium iodide and 2.6 g (3.9 mmol) (E)-3-[4-[[3-[(6-chlorohexyl)oxy]-5-[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]phenoxy]methyl]cyclohexyl]methoxy]-3-methoxyphenyl]acrylic acid methyl ester dissolved in 10 ml N,N-dimethylformamide the resulting mixture was stirred at 80 °C for 20 hours. The reaction mixture was poured into 50 ml of an ice cold 1 M HCl solution. The aqueous phase was extracted three times with 50 ml ethyl acetate. The organic phase was washed with 50 ml of a saturated sodium bicarbonate solution and repeatedly with water, dried over magnesium sulfate, filtered and concentrated by vacuum evaporation. Purification of the residue by column chromatography on 270 g

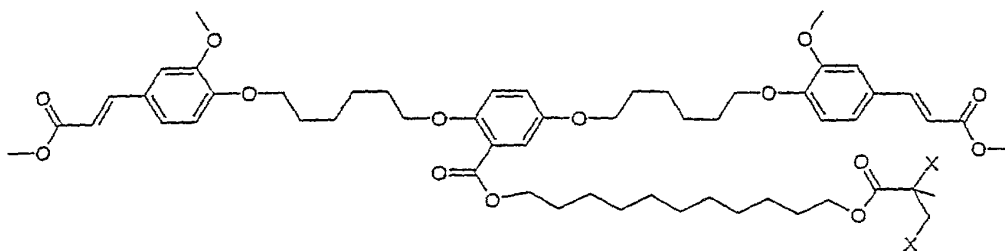
silica gel using toluene/ethanol (9:1) as eluent yielded 2.6 g (92%) of 2-methylacrylic acid

6-[3,5-bis[[2-methoxy-4-[(E)-2-(methoxycarbonyl)vinyl]-phenoxy]methyl]cyclohexyloxy]hexyl ester as yellowish oil.

5 Example 10

A two percent by weight solution S1 of the photoreactive polymer A was prepared using cyclopentanone as a solvent. The solution was stirred for 30 minutes at room temperature.

Photopolymer A:



15 Solution S1 was spin-coated at 2000 rpm onto two ITO (indium-tin-oxide) coated glass plates, which were then dried for 30 minutes at 150°C.

Both substrates were subsequently exposed for one minute to the polarised ultraviolet light of a 200 W high pressure mercury lamp. The intensity of the uv-light at the substrates position was measured as 1.1 mW/cm².

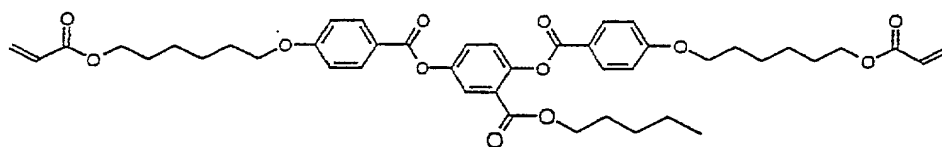
20 With the coated sides facing inwards, the two plates were assembled into a parallel liquid crystal cell which was filled with the nematic liquid crystal mixture MLC12000-000 (Merck). Using a polarising microscope, the alignment quality was found to be excellent. With a tilting compensator, which was introduced into the microscope the alignment of the long axis of the liquid crystal molecules was found to

be parallel to the polarisation direction of the uv-light which was used to photo-align photopolymer A.

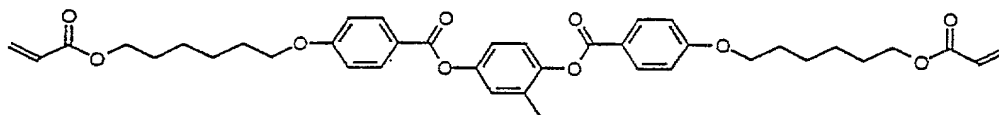
Example 11

5 A mixture M_{LCP} was prepared comprising the following liquid crystalline diacrylate monomers:

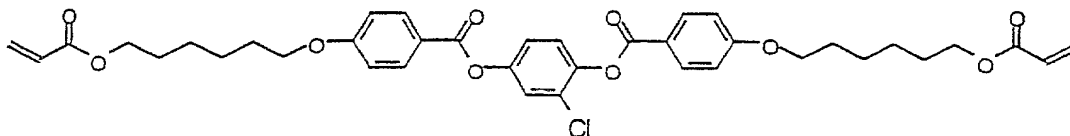
Mon1:



Mon2:



Mon3:



In addition to the diacrylate monomers, photoinitiator IRGACURE 369 from Ciba SC as well as BHT, which served as an inhibitor were added to the mixture. Thus the composition of mixture M_{LCP} was as follows:

Mon1	77 wt%
Mon2	14.5 wt%
Mon3	4.7 wt%
Irgacure 369	1.9 wt%
BHT	1.9 wt%

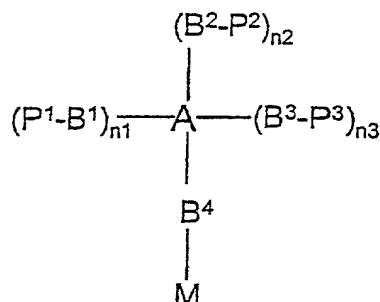
Finally, the solution $S(LCP)$ resulted from dissolving 15 wt% of mixture M_{LCP} in anisole.

Like in example 10, solution S1 was spincoated on a glass plate, dried and subsequently exposed to vertical incident, polarised uv-light for 1 minute. Then diacrylate solution $S(LCP)$ was spin-coated at 800 rpm for 2 minutes on top of the irradiated LPP-layer. To cross-link the liquid crystalline diacrylates the plate was exposed to isotropic uv-light under nitrogen atmosphere for 5 minutes.

To characterise the alignment capability of photopolymer A the contrast of the cross-linked liquid crystal layer was measured using a polarising microscope with crossed polarisers, which was further equipped with a photodiode for light intensity measurements. The contrast was calculated as the ratio of the light intensities measured with the optical axis of the cross-linked liquid crystal layer oriented 45° and 0° in respect to one of the polarisers. The high contrast of 1200:1 demonstrates the excellent alignment capability of photopolymer A.

Claims

1. A compound comprising a repeating unit of formula (I)



I

in which:

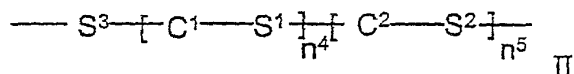
A represents a nitrogen atom, a carbon atom, a group $-CR^1-$ or an aromatic or alicyclic group, which is optionally substituted by a group selected from fluorine, chlorine, cyano and a C_{1-18} cyclic, straight-chain or branched alkyl group, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkyl $-CH_2-$ groups are optionally replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-Si(CH_3)_2-O-Si(CH_3)_2-$, $-NR^1-$, $-NR^1-CO-$, $-CO-NR^1-$, $-NR^1-CO-O-$, $-O-CO-NR^1-$, $-NR^1-CO-NR^1-$, $-CH=CH-$, $-C\equiv C-$ and $-O-CO-O-$, wherein R^1 represents a hydrogen atom or lower alkyl,

M represents a repeating monomer unit;

n^1 to n^3 each independently represent 0 or an integer having a value of from 1 to 3, with the proviso that $1 < n^1 + n^2 + n^3 < 4$;

P^1, P^2, P^3 each independently represents a photoactive group; and

B¹ to B⁴ each independently represent a residue of general formula II



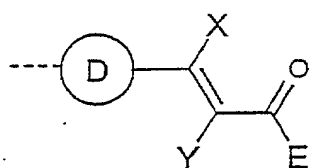
in which

5 S¹ to S³ each independently represent a single bond or a spacer group selected from a C₁₋₂₄ straight-chain or branched alkylene group, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkylene -CH₂- groups are optionally replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-,
10 -Si(CH₃)₂-O-Si(CH₃)₂-, -NR¹-, -NR¹-CO-, -CO-NR¹-,
-NR¹-CO-O-, -O-CO-NR¹-, -NR¹-CO-NR¹-, -CH=CH-, -C≡C- and -O-CO-O- wherein R¹ is as defined above,

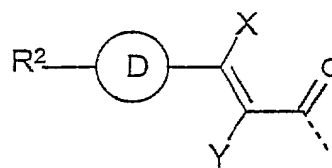
C¹ and C² each independently represents an aromatic or an alicyclic group, which is optionally substituted by a group selected from fluorine, chlorine, cyano or a C₁₋₁₈ cyclic, straight-chain or branched alkyl group, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkyl -CH₂- groups are optionally replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-,
15 -Si(CH₃)₂-O-Si(CH₃)₂-, -NR¹-, -NR¹-CO-, -CO-NR¹-,
20 -NR¹-CO-O-, -O-CO-NR¹-, -NR¹-CO-NR¹-, -CH=CH-, -C≡C- and -O-CO-O- wherein R¹ represents a hydrogen atom or lower alkyl, and

n⁴ and n⁵ are each independently 0 or 1.

2. A compound according to Claim 1, in which P¹ to P³ are selected from the general
25 formulae IIIa and IIIb:



IIIa



IIIb

5 wherein the broken line indicates the point of linkage to S^3 and wherein:

D represents pyrimidine-2,5-diyl, pyridine-2,5-diyl, 2,5-thiophenylene, 2,5-furanylene, 1,4- or 2,6-naphthylene; a phenylene group, which is optionally substituted by a group selected from fluorine, chlorine, cyano; or a C_{1-18} cyclic, straight-chain or branched alkyl residue, which is optionally substituted by a single cyano group or by one or more halogen groups and in which one or more non-adjacent alkyl $-CH_2-$ groups are optionally replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-Si(CH_3)_2-O-Si(CH_3)_2-$, $-NR^1-$, $-NR^1-CO-$, $-CO-NR^1-$, $-NR^1-CO-O-$, $-O-CO-NR^1-$, $-NR^1-CO-NR^1-$, $-CH=CH-$, $-C\equiv C-$ and $-O-CO-O-$, wherein R^1 is as defined above;

E represents $-OR^3$, $-NR^4R^5$ or an oxygen atom, which defines together with the ring D a coumarin unit, wherein R^3 , R^4 and R^5 are selected from hydrogen and a C_{1-18} cyclic, straight-chain or branched alkyl residue, which is optionally substituted by one or more halogen atoms and in which one or more non-adjacent alkyl $-CH_2-$ groups are optionally replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$ and $-CH=CH-$, or R^4 and R^5 together form a C_{5-8} alicyclic ring;

X, Y each independently represent hydrogen, fluorine, chlorine, cyano or a C_{1-12} alkyl group, which is optionally substituted by fluorine and in which one or

more non-adjacent alkyl $-\text{CH}_2-$ groups are optionally replaced by a group selected from $-\text{O}-$, $-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-$ and $-\text{CH}=\text{CH}-$;

R^2 represents hydrogen or a C_{1-18} straight-chain or branched alkyl residue, which is optionally substituted by a single cyano group or by one or more halogen atoms and in which one or more non-adjacent alkyl $-\text{CH}_2-$ groups are independently optionally replaced by a group selected from $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-$, $-\text{Si}(\text{CH}_3)_2-\text{O}-\text{Si}(\text{CH}_3)_2-$, $-\text{NR}^1-$, $-\text{NR}^1-\text{CO}-$, $-\text{CO}-\text{NR}^1-$, $-\text{NR}^1-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-\text{NR}^1-$, $-\text{NR}^1-\text{CO}-\text{NR}^1-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$ and $-\text{O}-\text{CO}-\text{O}-$, wherein R^1 is as defined above.

3. A compound according to Claim 1 or Claim 2, in which the repeating unit of formula (I) comprises at least 50% of the monomer building blocks comprising the compound of formula (I).
4. A compound according to any one of claims 1 to 3, in which the group M is selected from acrylate; methacrylate; 2-chloroacrylate; 2-phenylacrylate; acrylamide, methacrylamide, 2-chloroacrylamide and 2-phenylacrylamide, the nitrogen atom of which is optionally substituted by a lower alkyl group; vinyl ether; vinyl ester; a styrene derivative; siloxane; imide; amic acid; amic acid esters; amidimide; maleic acid derivatives and fumaric acid derivatives.
5. A method of manufacturing a compound of formula (I) comprising the polymerisation of one or more pre-finished monomer units of formula (I).
6. A method of manufacturing a compound of formula (I) comprising reacting a photoactive derivative with a functional polymer analogue of a polymer according to Claim 1.
7. A polymer layer comprising a compound of formula (I) in cross-linked form.
8. A polymer layer according to Claim 7, which is an orientation layer for an optical or an electro-optical device.

9. Use of a compound according to any one of claims 1 to 4 in the manufacture of an optical or an electro-optical device.

10. An optical or an electro-optical device comprising a compound according to any one of claims 1 to 4.

5 11. An optical or an electro-optical device comprising a layer according to Claim 7 or Claim 8.

12. A compound of formula (I), which is Poly-[1-[11-[5-[4-[(*E*)-2-methoxycarbonylvinyl]benzoyloxy]-2-[6-[2-methoxy-(*E*)-4-(methoxycarbonylvinyl)-phenoxy]oxyhexyl]benzoyloxy]undecyloxy]carbonyl]-1-methylethylene]

10 13. A compound of formula (I), which is Poly-[1-[11-[(*E,E*)-2,5-di-[6-[2-methoxy-4-(methoxycarbonylvinyl)phenoxy]oxyhexyl]benzoyloxy]undecyloxy]carbonyl]-1-methylethylene].

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that: My residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Compound

the specification of which:

is attached hereto; or

was filed as United States Application Serial No. **serial number**
on **filing date**, and was amended on _____
(if applicable); or

was filed as PCT International Application Number PCT/EP00/06788
on July 17, 2000.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate or § 365(a) of any PCT international application(s), designating at least one country other than the United States, listed below and have also identified below any foreign application(s) for patent or inventor's certificate, or any PCT international application(s) having a filing date before that of the application(s) of which priority is claimed:

Country	Application Number	Date of Filing	Priority Claimed Under 35 U.S.C. 119
Europe	EP 99 305 857.7	July 23, 1999	<input checked="" type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO

I hereby claim the benefit under 35 U.S.C. § 119(e) of any United States provisional application(s) listed below:

Application Number	Date of Filing

I hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s) or § 365(c) of any PCT international application(s) designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT international application(s) in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application(s) and the national or PCT international filing date of this application:

Application Number	Date of Filing	Status (Patented, Pending, Abandoned)

I hereby appoint the following attorney and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith. **FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, L.L.P., Reg. No. 22,540**, Douglas B. Henderson, Reg. No. 20,291; Ford F. Farabow, Jr., Reg. No. 20,630; Arthur S. Garrett, Reg. No. 20,338; Donald R. Dunner, Reg. No. 19,073; Brian G. Brunsvold, Reg. No. 22,593; Tipton D. Jennings IV, Reg. No. 20,645; Jerry D. Voight, Reg. No. 23,020; Laurence R. Heffer, Reg. No. 20,827; Kenneth E. Payne, Reg. No. 23,098; Herbert H. Mintz, Reg. No. 26,691; C. Larry O'Rourke, Reg. No. 26,014; Albert J. Santorelli, Reg. No. 22,610; Michael C. Elmer, Reg. No. 25,857; Richard H. Smith, Reg. No. 20,609; Stephen L. Peterson, Reg. No. 26,325; John M. Romary, Reg. No. 26,331; Bruce C. Zotter, Reg. No. 27,680; Dennis P. O'Reilly, Reg. No. 27,932; Allen M. Sokal, Reg. No. 26,695; Robert D. Bajefsky, Reg. No. 25,387; Richard L. Stroup, Reg. No. 28,478; David W. Hill, Reg. No. 28,220; Thomas L. Irving, Reg. No. 28,619; Charles E. Lipsey, Reg. No. 28,165; Thomas W. Winland, Reg. No. 27,605; Basil J. Lewis, Reg. No. 28,818; Martin I. Fuchs, Reg. No. 28,508; E. Robert Yoches, Reg. No. 30,120; Barry W. Graham, Reg. No. 29,924; Susan Haberman Griffin, Reg. No. 30,907; Richard B. Racine, Reg. No. 30,415; Thomas H. Jenkins, Reg. No. 30,857; Robert E. Converse, Jr., Reg. No. 27,432; Clair X. Mullen, Jr., Reg. No. 20,348; Christopher P. Foley, Reg. No. 31,354; John C. Paul, Reg. No. 30,413; David M. Kelly, Reg. No. 30,953; Kenneth J. Meyers, Reg. No. 25,146; Carol P. Einaudi, Reg. No. 32,220; Walter Y. Boyd, Jr., Reg. No. 31,738; Steven M. Anzalone, Reg. No. 32,095; Jean B. Fordis, Reg. No. 32,984; Barbara C. McCurdy, Reg. No. 32,120; James K. Hammond, Reg. No. 31,964; Richard V. Burgujian, Reg. No. 31,744; J. Michael Jakes, Reg. No. 32,824; Dirk D. Thomas, Reg. No. 32,600; Thomas W. Banks, Reg. No. 32,719; Christopher P. Isaac, Reg. No. 32,616; Bryan C. Diner, Reg. No. 32,409; M. Paul Barker, Reg. No. 32,013; Andrew Chanho Sonu, Reg. No. 33,457; David S. Forman, Reg. No. 33,694; Vincent P. Kovalick, Reg. No. 32,867; James W. Edmondson, Reg. No. 33,871; Michael R. McGurk, Reg. No. 32,045; Joann M. Neth, Reg. No. 36,363; Gerson S. Panitch, Reg. No. 33,751; Cheri M. Taylor, Reg. No. 33,216; Charles E. Van Horn, Reg. No. 40,266; and Linda A. Wadler, Reg. No. 33,218; Jeffrey A. Berkowitz, Reg. No. 36,743; Michael R. Kelly, Reg. No. 33,921; James B. Monroe, Reg. No. 33,971; Doris Johnson Hines, Reg. No. 34,629; Allen R. Jensen, Reg. No. 28,224; Lori Ann Johnson, Reg. No. 34,498; and David A. Manspeizer, Reg. No. 37,540 and Steven J. Scott, Reg. No. 43,911. Please address all correspondence to **FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, L.L.P., 1300 I Street, N.W., Washington, D.C. 20005, Telephone No. (202) 408-4000.**

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

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